



US009409975B2

(12) **United States Patent**
Hostetter et al.

(10) **Patent No.:** **US 9,409,975 B2**
(45) **Date of Patent:** **Aug. 9, 2016**

(54) **ANTIBODY BINDING MICROBIAL HEPARIN BINDING MOTIF TO RETARD OR PREVENT MICROBIAL BIOFILM FORMATION ON IMPLANTED MEDICAL DEVICES**

(58) **Field of Classification Search**
None
See application file for complete search history.

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(57) **ABSTRACT**

Methods and reagents for ameliorating biofilm formation on a surface of an indwelling or implanted device in a patient resulting in decreased virulence of microorganisms such as *Candida* species and/or *Staphylococcus* species.

8 Claims, 21 Drawing Sheets

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(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 0 days.

(21) Appl. No.: **14/518,051**

(22) Filed: **Oct. 20, 2014**

(65) **Prior Publication Data**

US 2015/0050284 A1 Feb. 19, 2015

Related U.S. Application Data

(63) Continuation-in-part of application No. PCT/US2013/031499, filed on Mar. 14, 2013.

(60) Provisional application No. 61/636,243, filed on Apr. 20, 2012.

(51) **Int. Cl.**

A61K 31/00 (2006.01)

A61K 39/395 (2006.01)

C07K 16/00 (2006.01)

C07K 16/14 (2006.01)

C07K 16/12 (2006.01)

C07K 7/08 (2006.01)

A61K 39/00 (2006.01)

(52) **U.S. Cl.**

CPC . **C07K 16/14** (2013.01); **C07K 7/08** (2013.01); **C07K 16/1271** (2013.01); **A61K 39/00** (2013.01); **A61K 2039/505** (2013.01); **A61K 2039/575** (2013.01); **C07K 2317/34** (2013.01); **C07K 2317/76** (2013.01)

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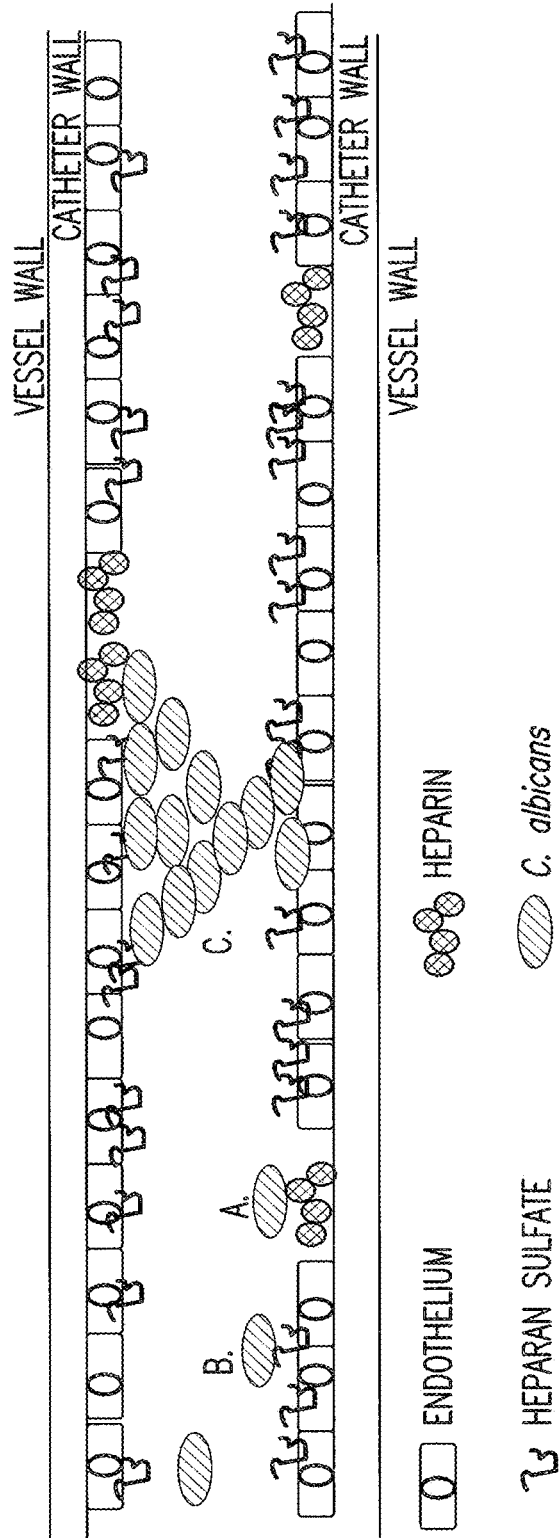


FIG.1

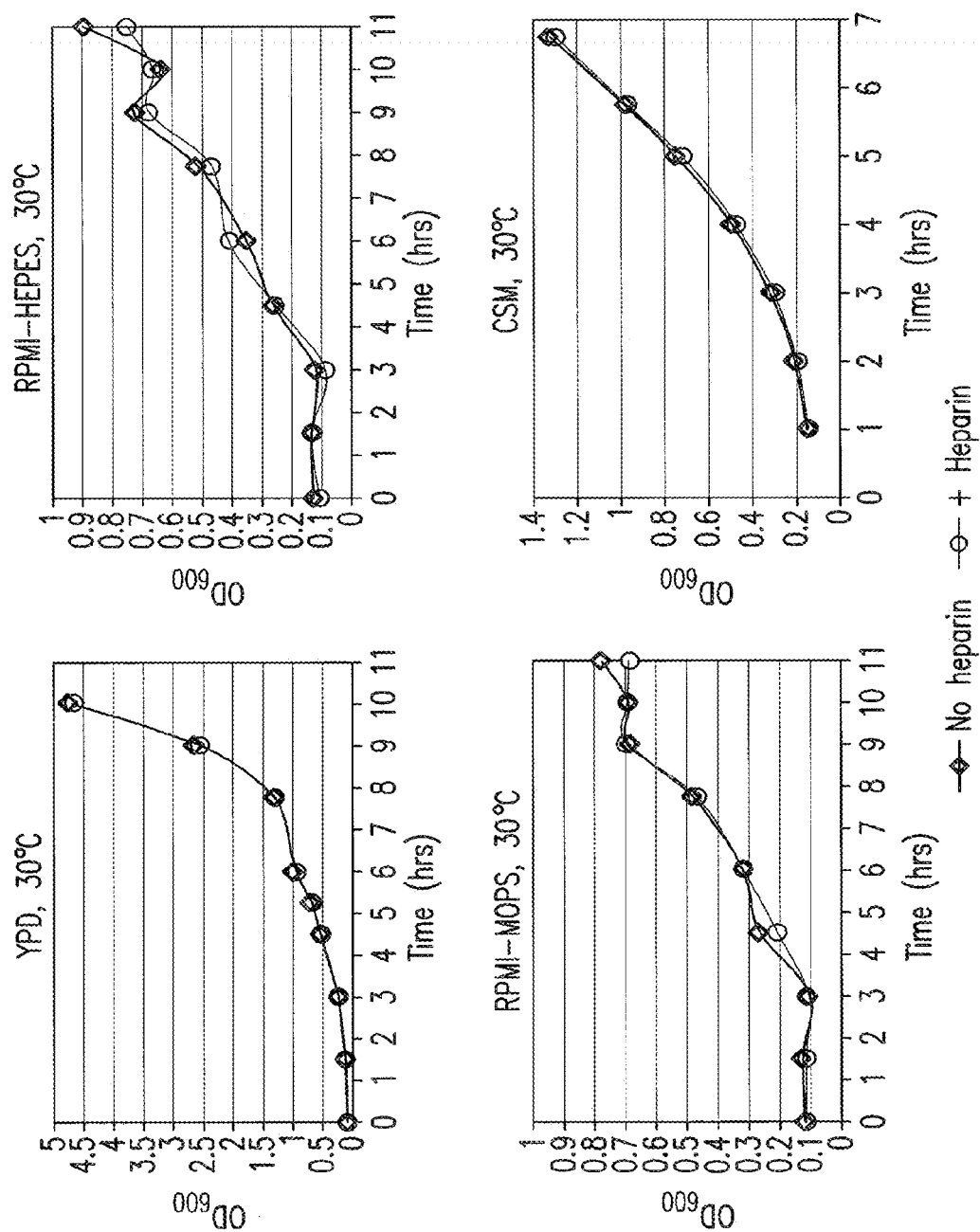


FIG.2A

Doubling Times (hours) at 30°C		
	No Heparin	+ Heparin
YPD	1.70	1.69
RPMI-HEPES	3.56	3.75
RPMI-MOPS	3.58	3.51
CSM	1.82	1.79

FIG.2B

Percent Hyphae		
	No Heparin	+ Heparin
YPD, 30 °C, 2.5 hours	0.13	0.09
RPMI-HEPES, 37 °C, 1 hour	7.37	5.01

FIG.2C

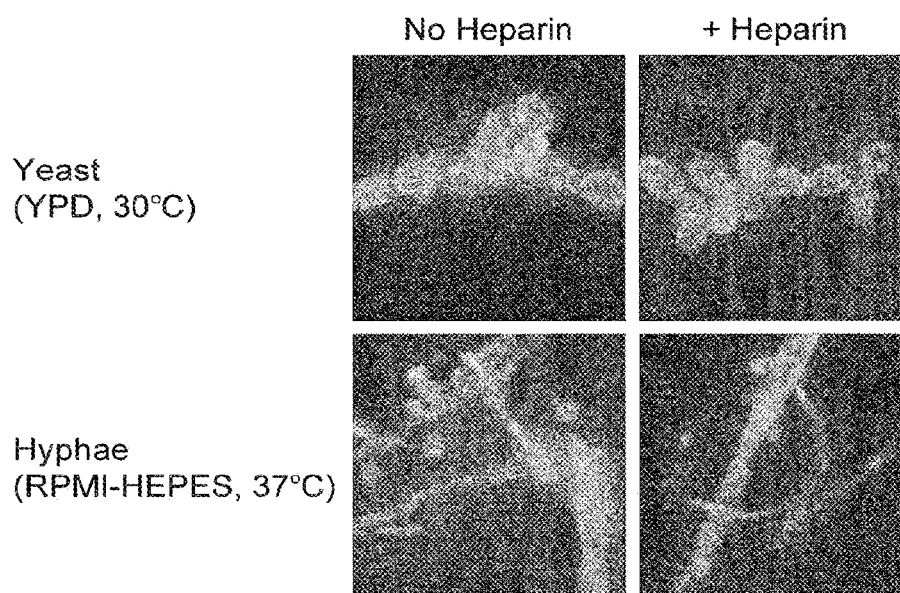


FIG.2D

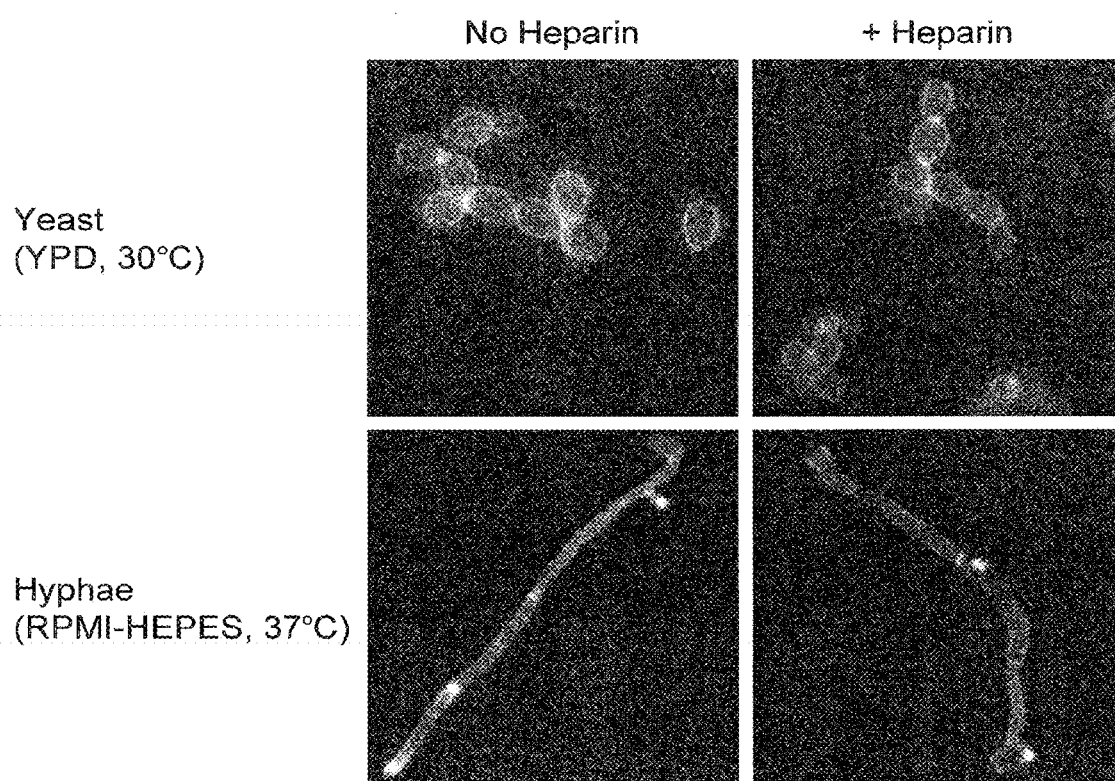


FIG.2E

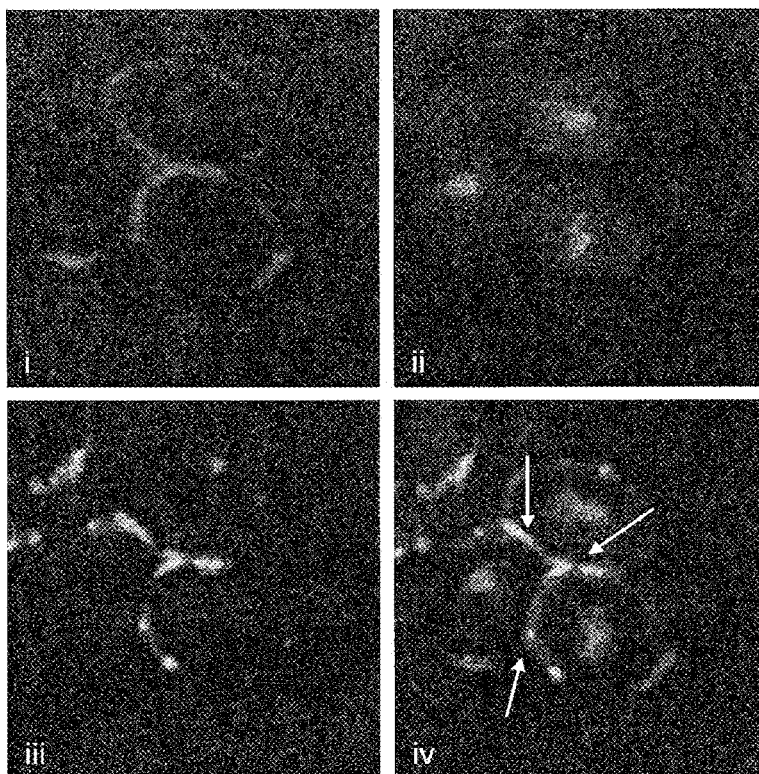


FIG.3A

Optical Density ₅₉₅ at 10,000 CFU/well <i>C. albicans</i>					
Heparin units/well ($\mu\text{g}/\text{well}$)	1.25 (6.3)	2.5 (12.5)	5.0 (25.0)	10.0 (50.0)	20.0 (100.0)
Mean OD ₅₉₅ (\pm SD)	0.53 (\pm 0.05)	0.51 (\pm 0.03)	0.48 (\pm 0.04)	0.51 (\pm 0.02)	0.49 (\pm 0.03)

FIG.3B

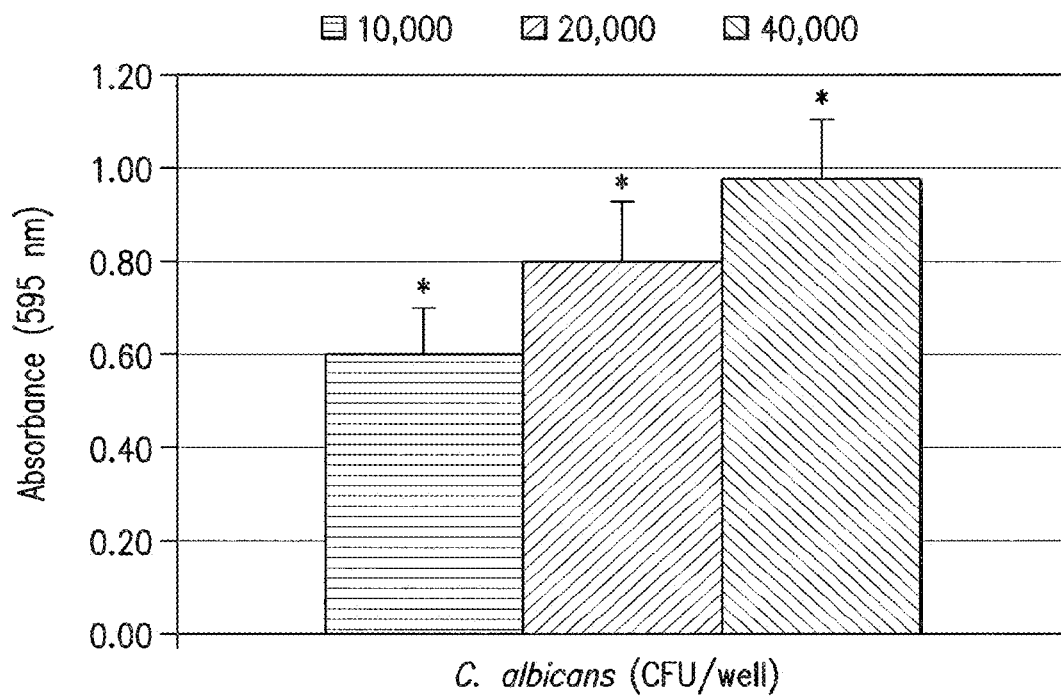


FIG.3C

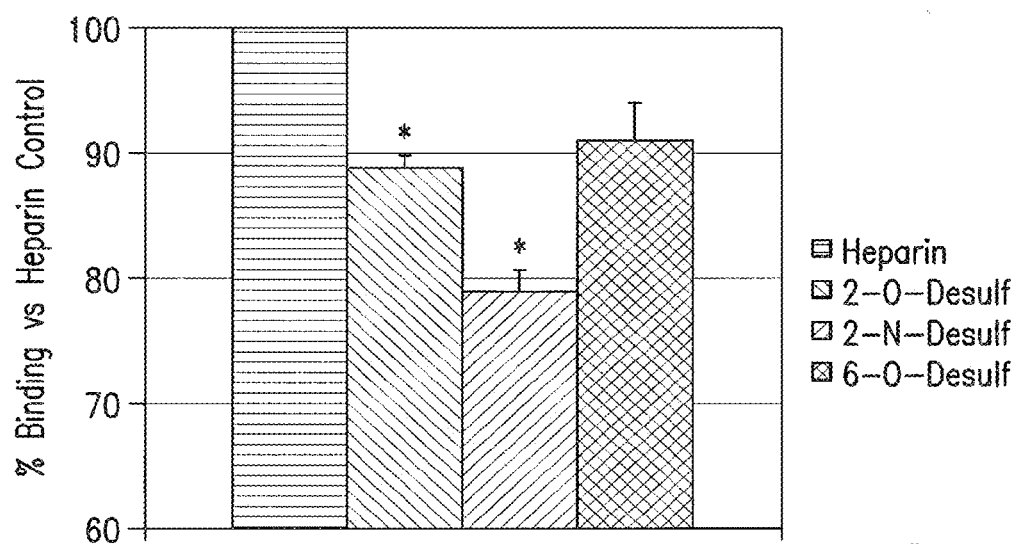


FIG.3D

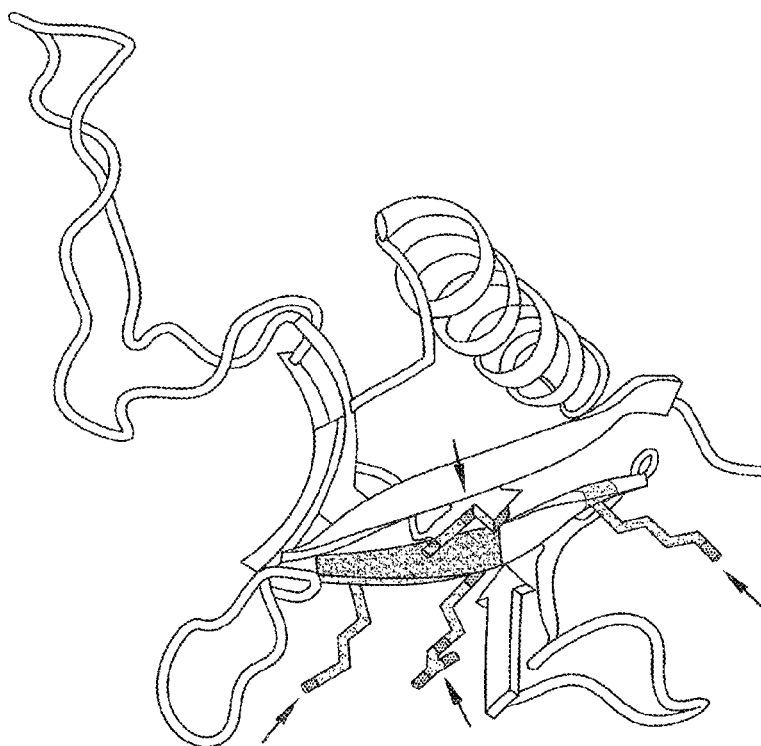
N = 3
*p ≤ 0.003

Motif 1	Motif 2	Motif 3	Motif 4	Motif 5
804 QKKHQIHK	1383 THKGRF	1530 MKRGKP	1593 FKKRFFKL	1612 SHKTRA

↓
804 QAAHQIHK

↓
1593 FKAAFFKL

FIG.4A



1593 FKKRFFKL
(Motif 4)

FIG.4B

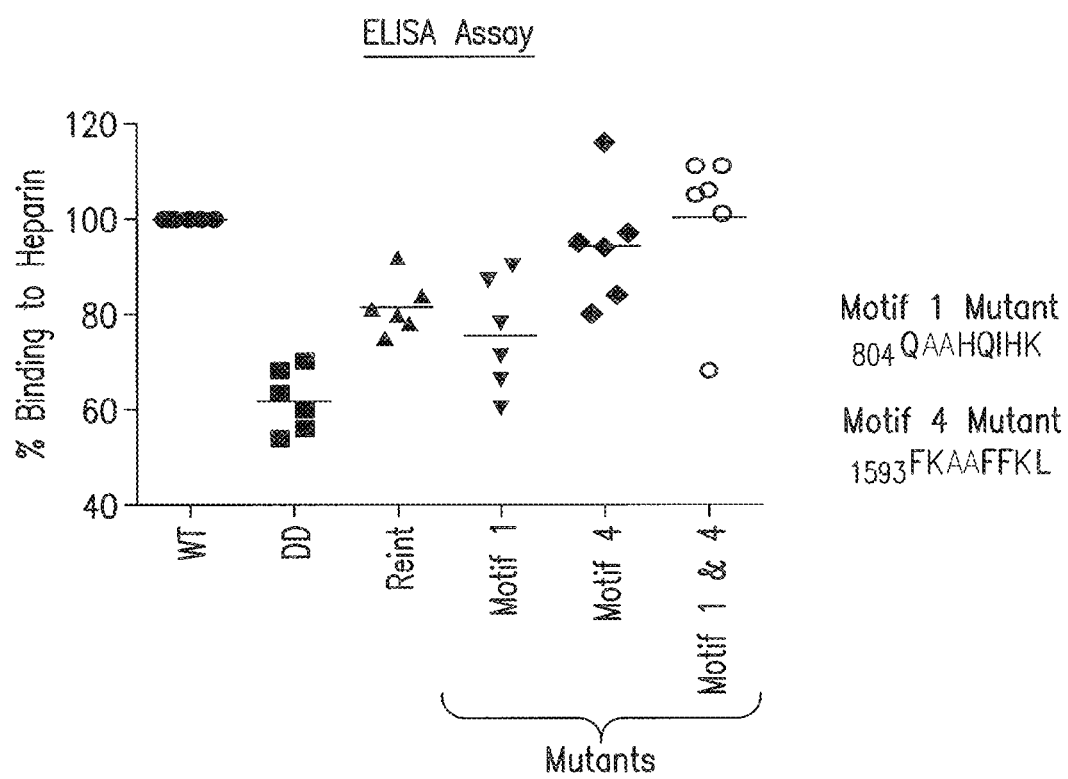


FIG.4C

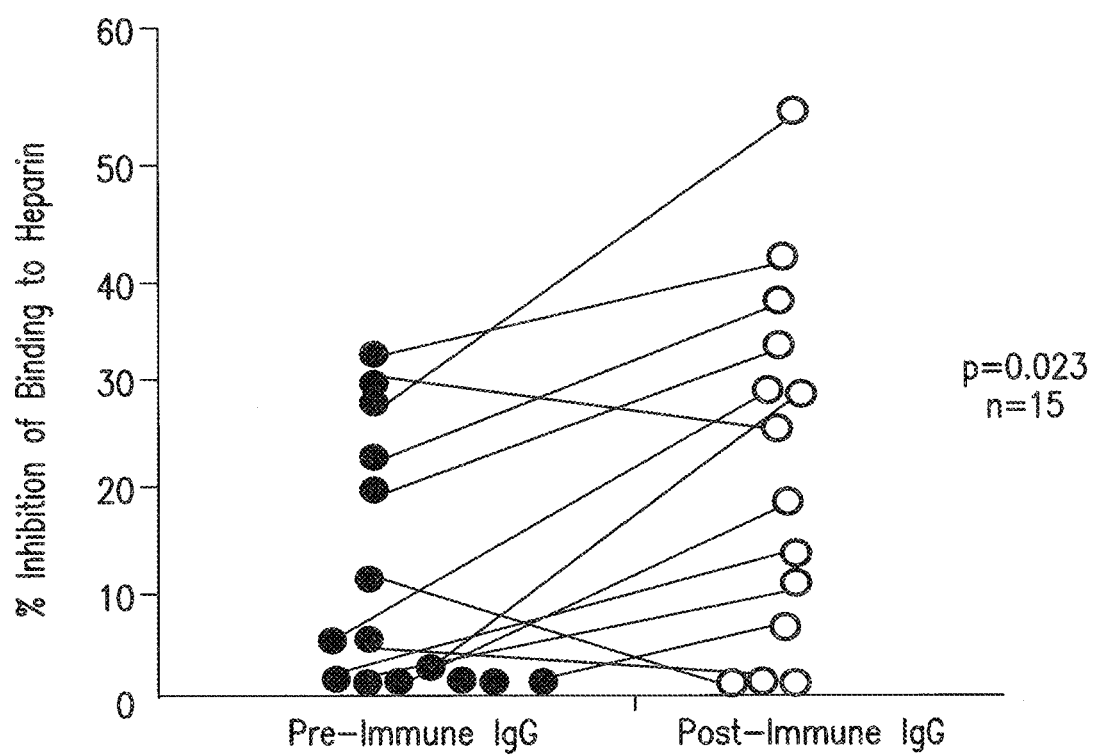


FIG. 5

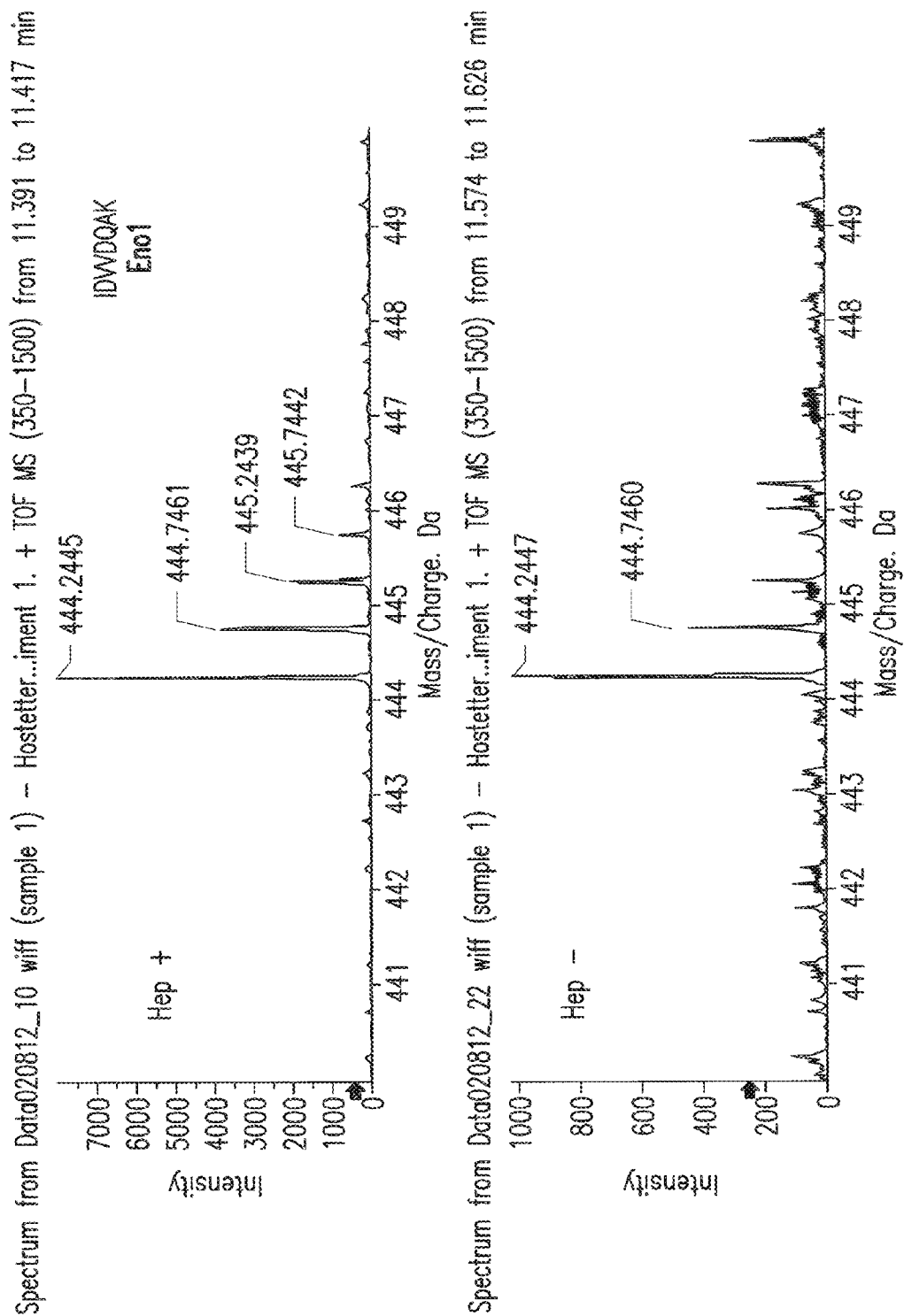


FIG. 6A

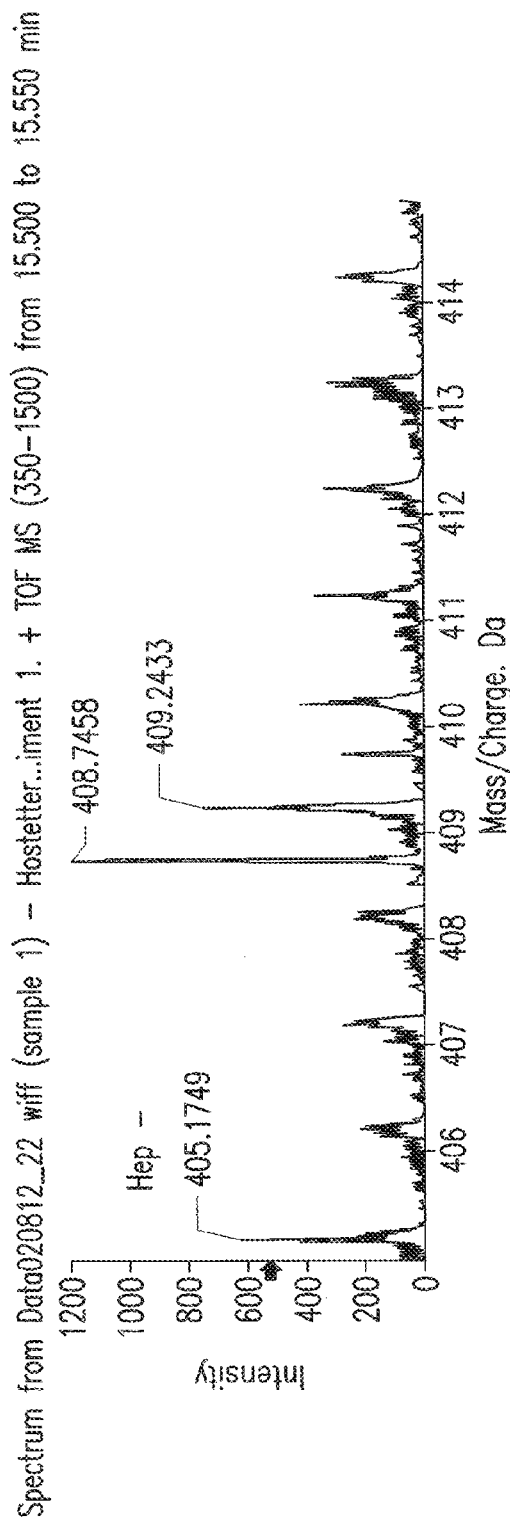
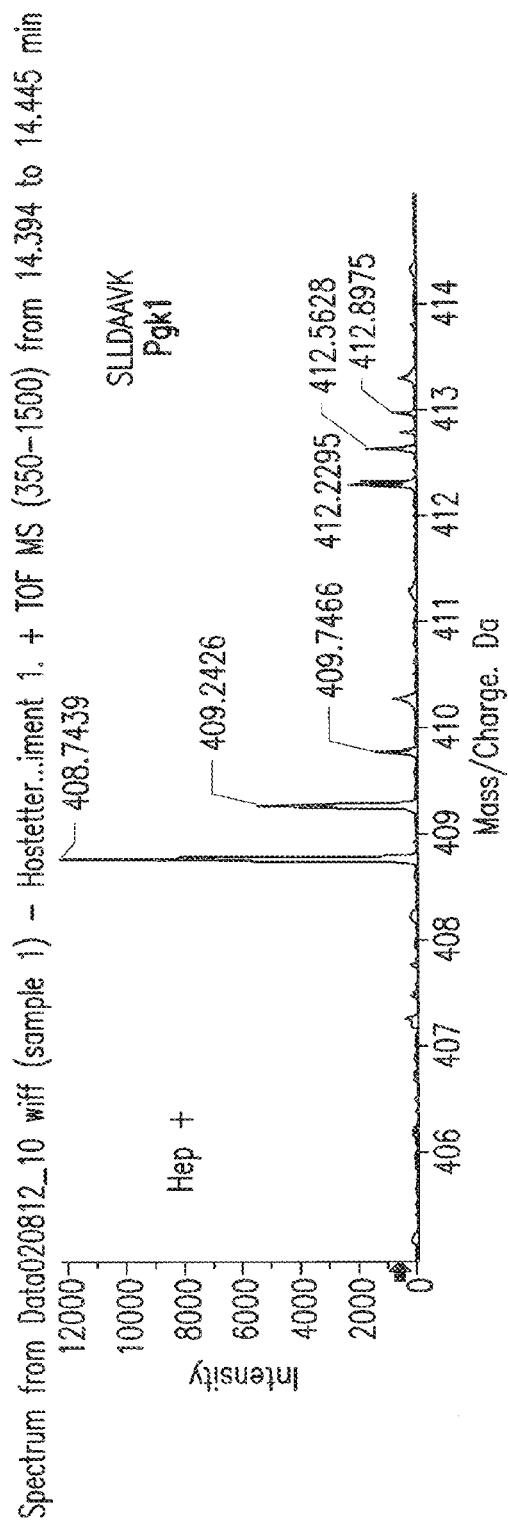


FIG. 6B

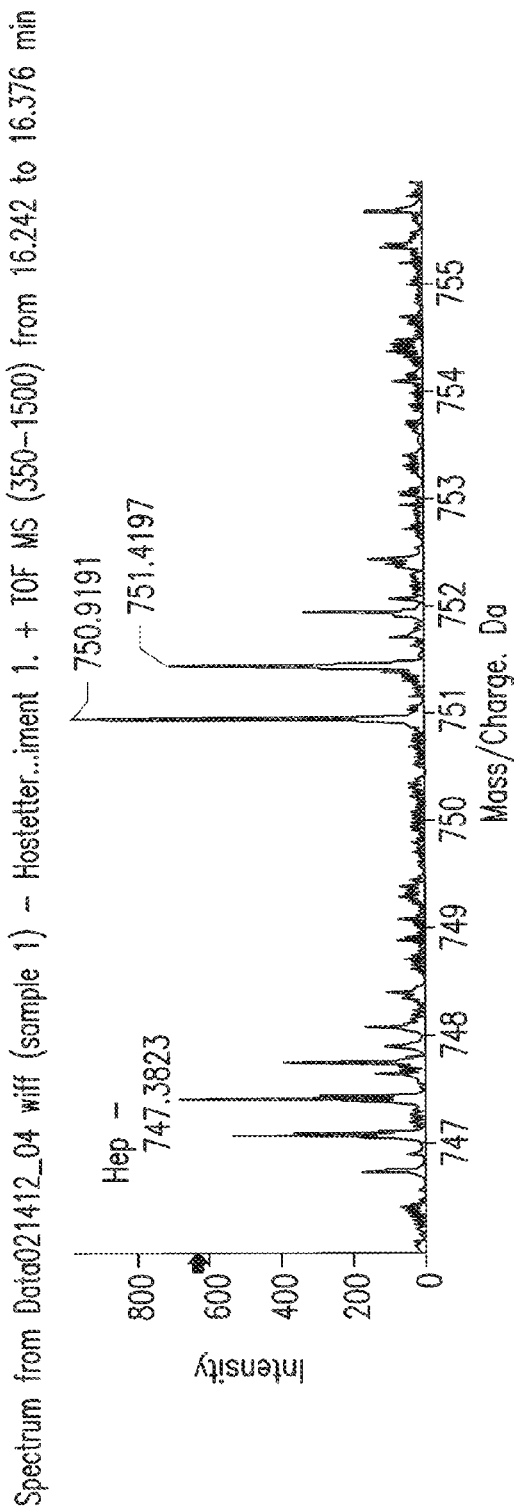
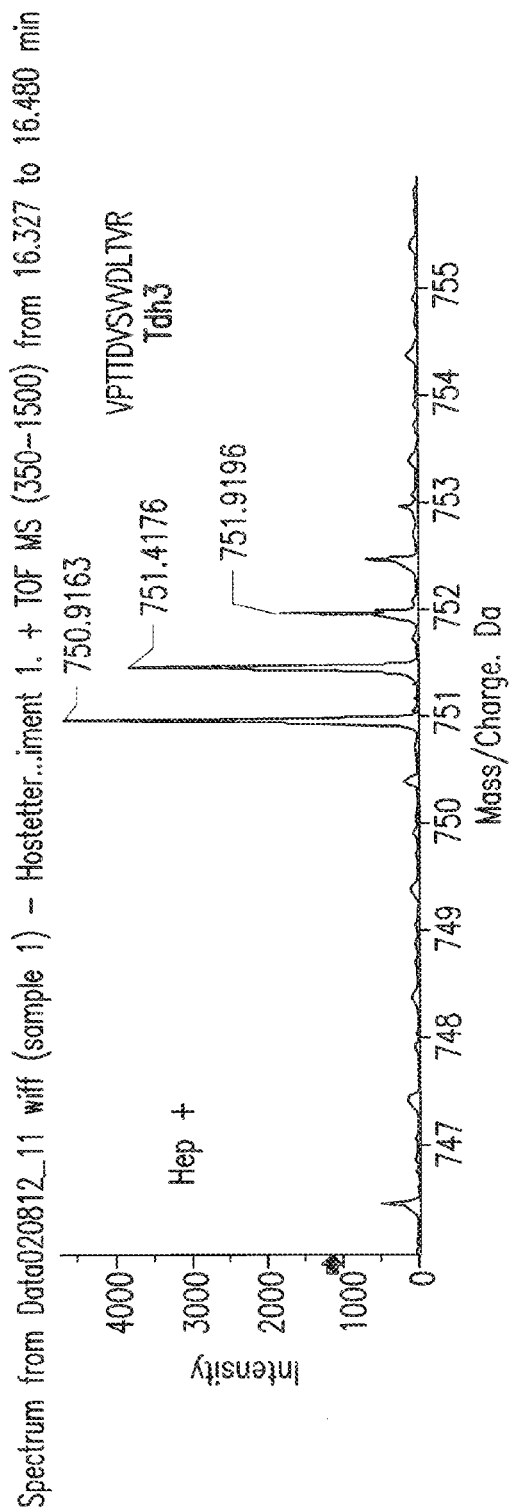


FIG. 6C

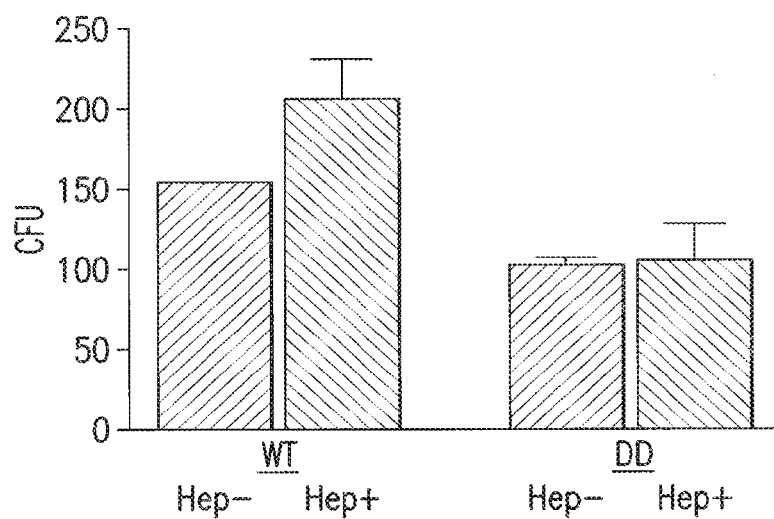


FIG. 7A

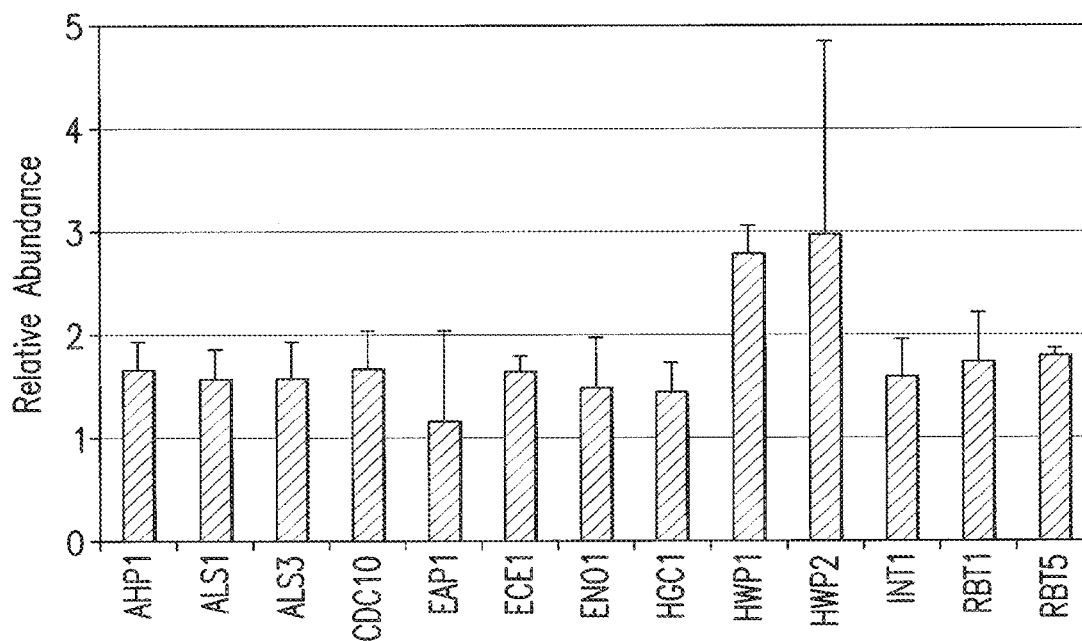


FIG. 7B

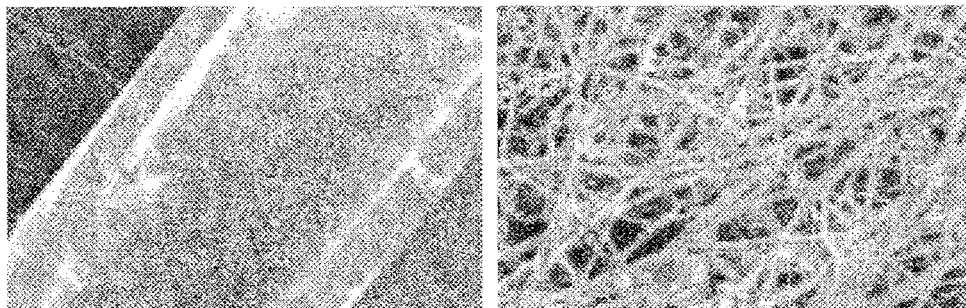
INT1 Wild Type (WT)

FIG.8A

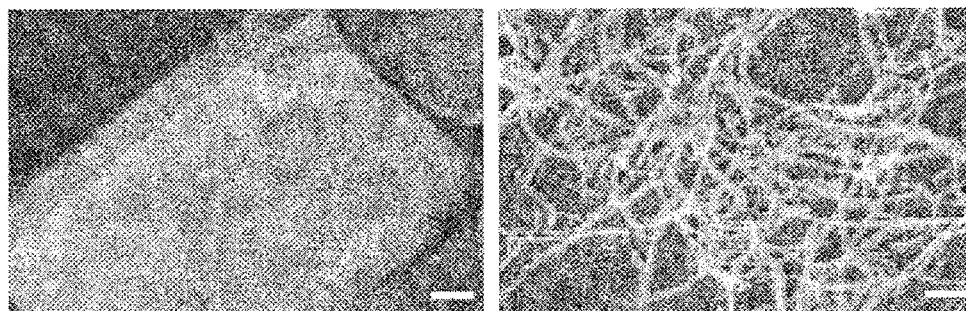
INT1 Double Disruptant (DD)

FIG.8B

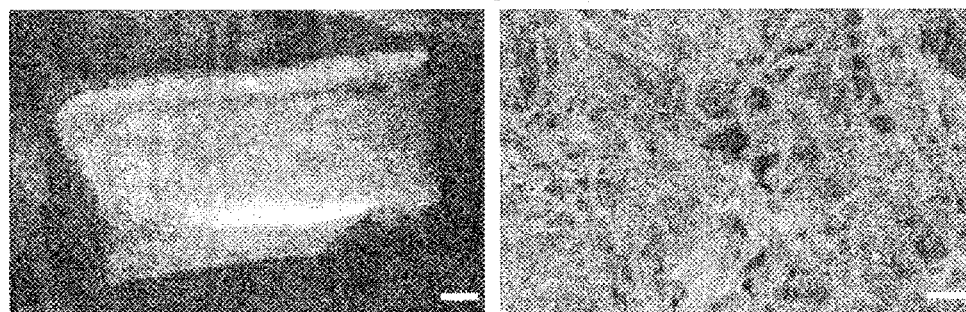
INT1 Reintegrant (Reint)

FIG.8C

Motif 1 Mutant

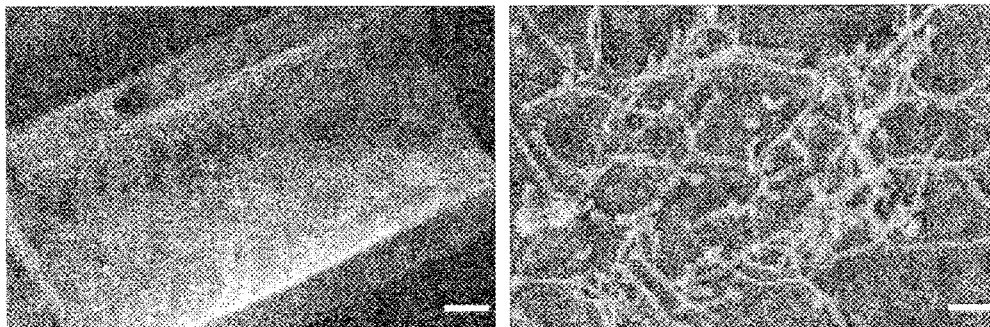


FIG.8D

Motif 4 Mutant

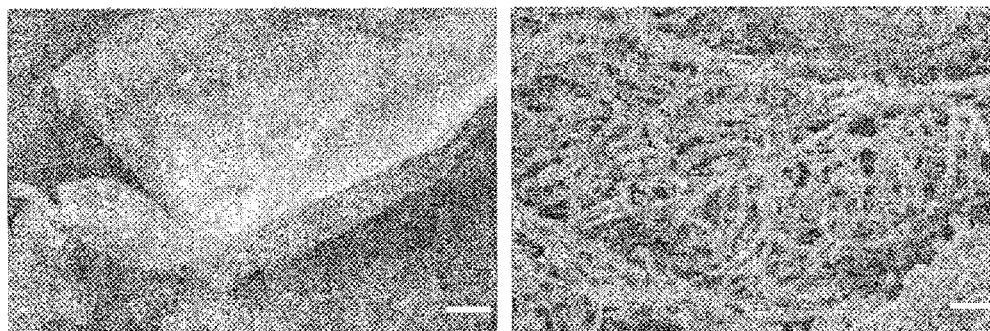


FIG.8E

Motif 1 & 4 Mutant

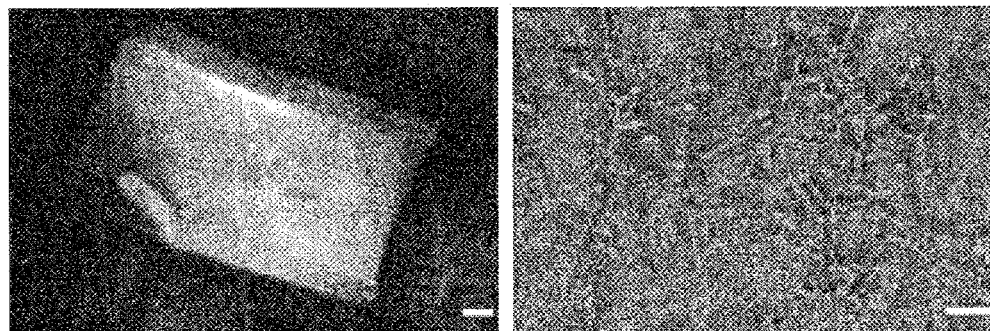


FIG.8F

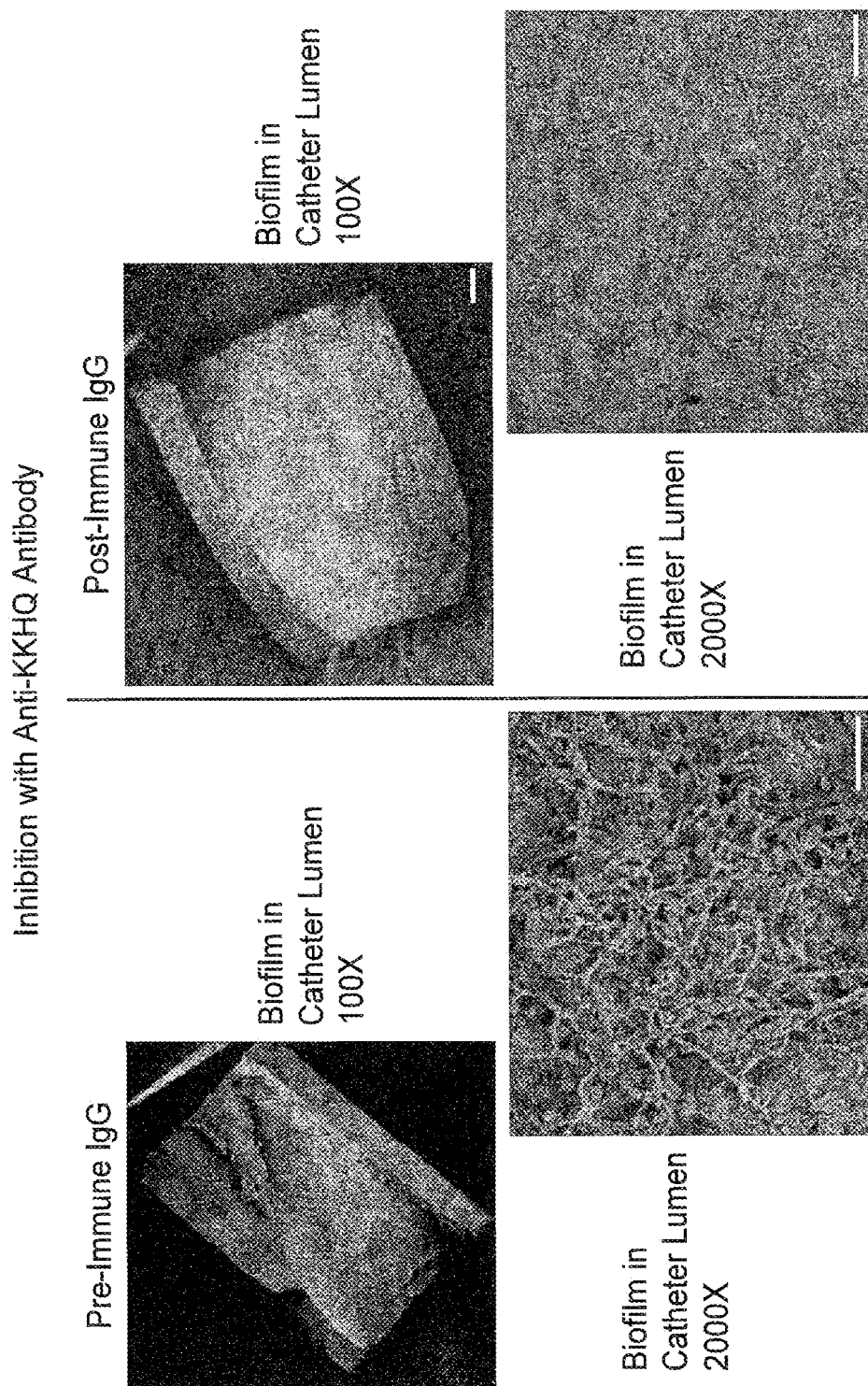


FIG.9

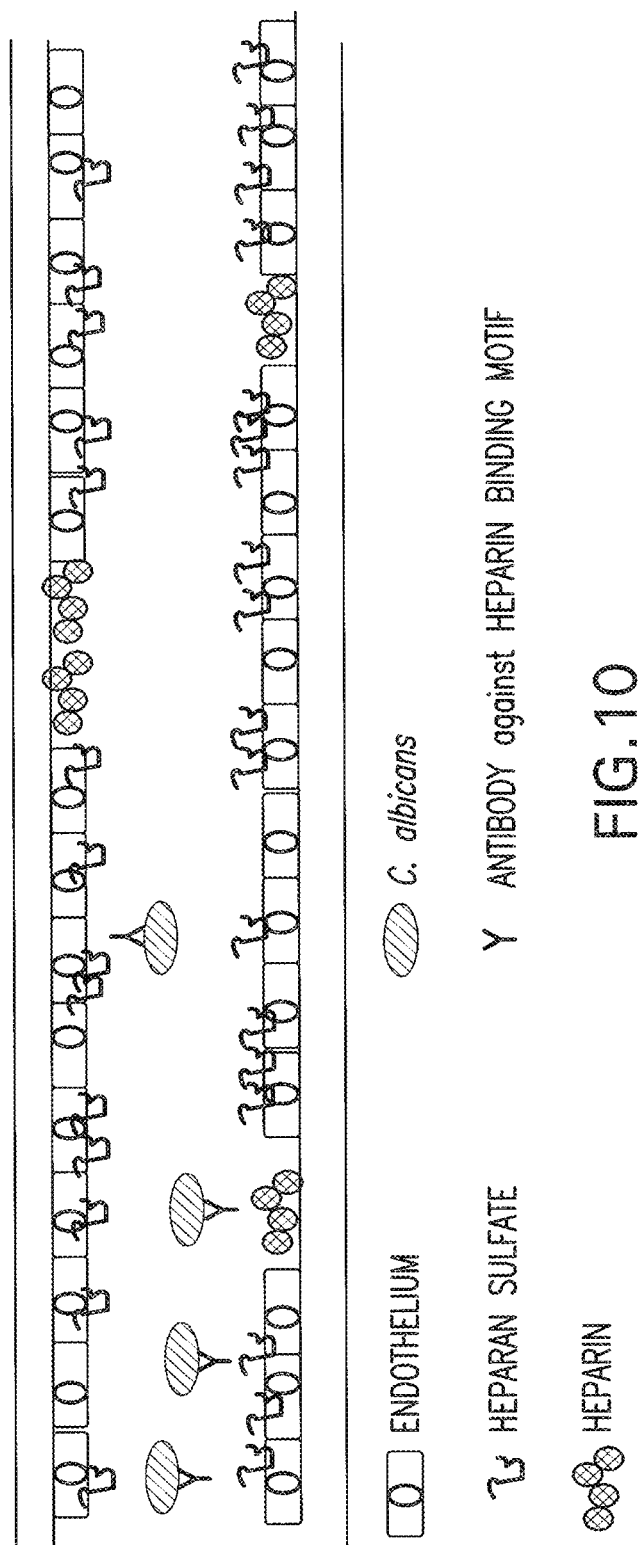


FIG. 11

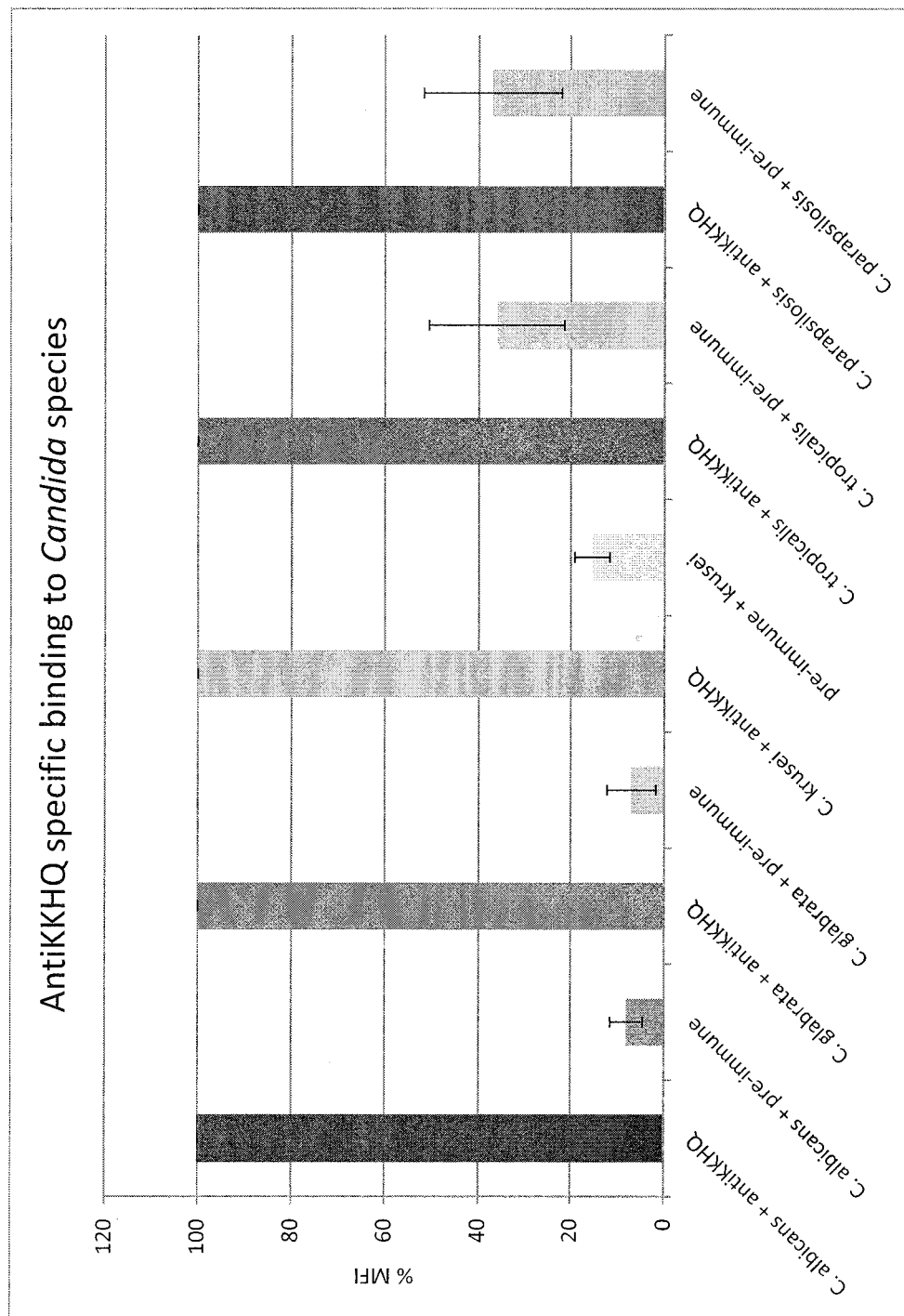


FIG. 12A

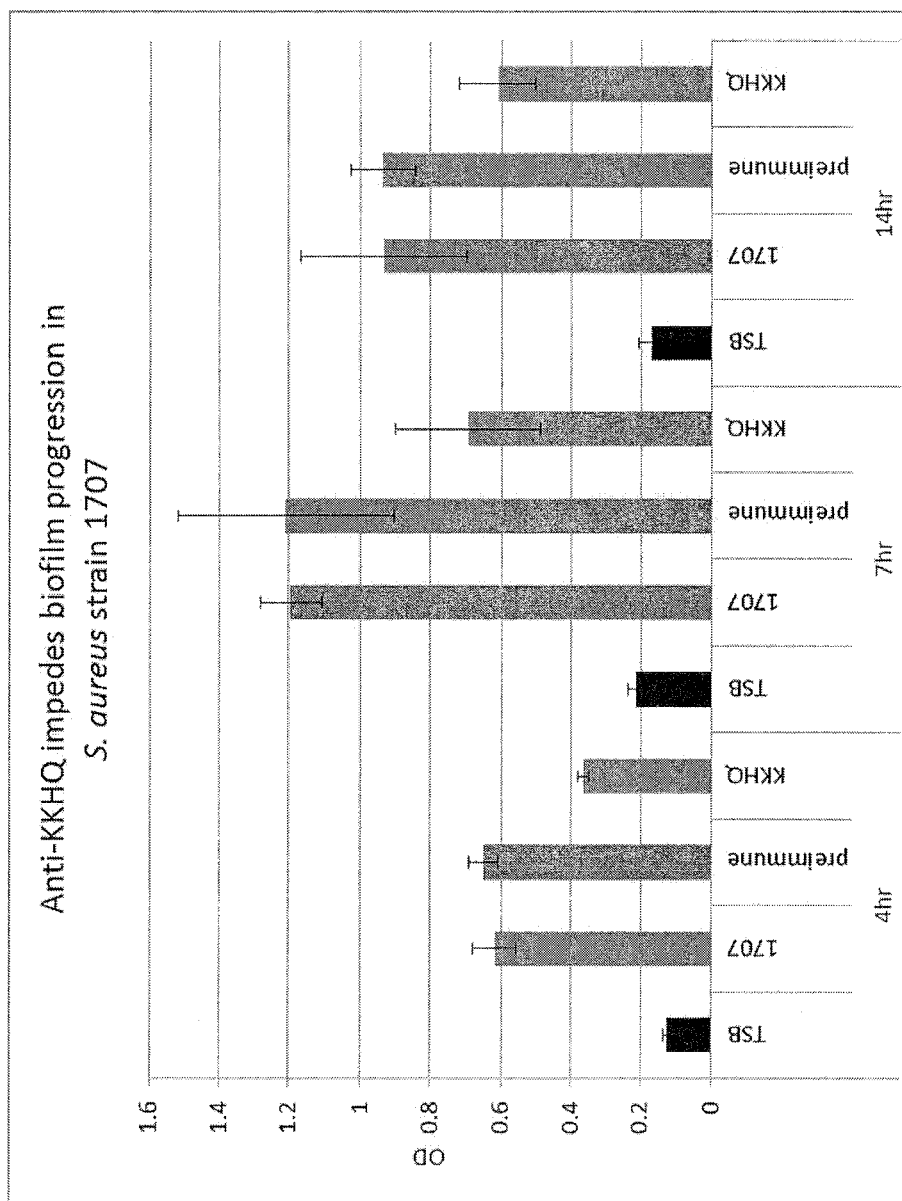


FIG. 12B

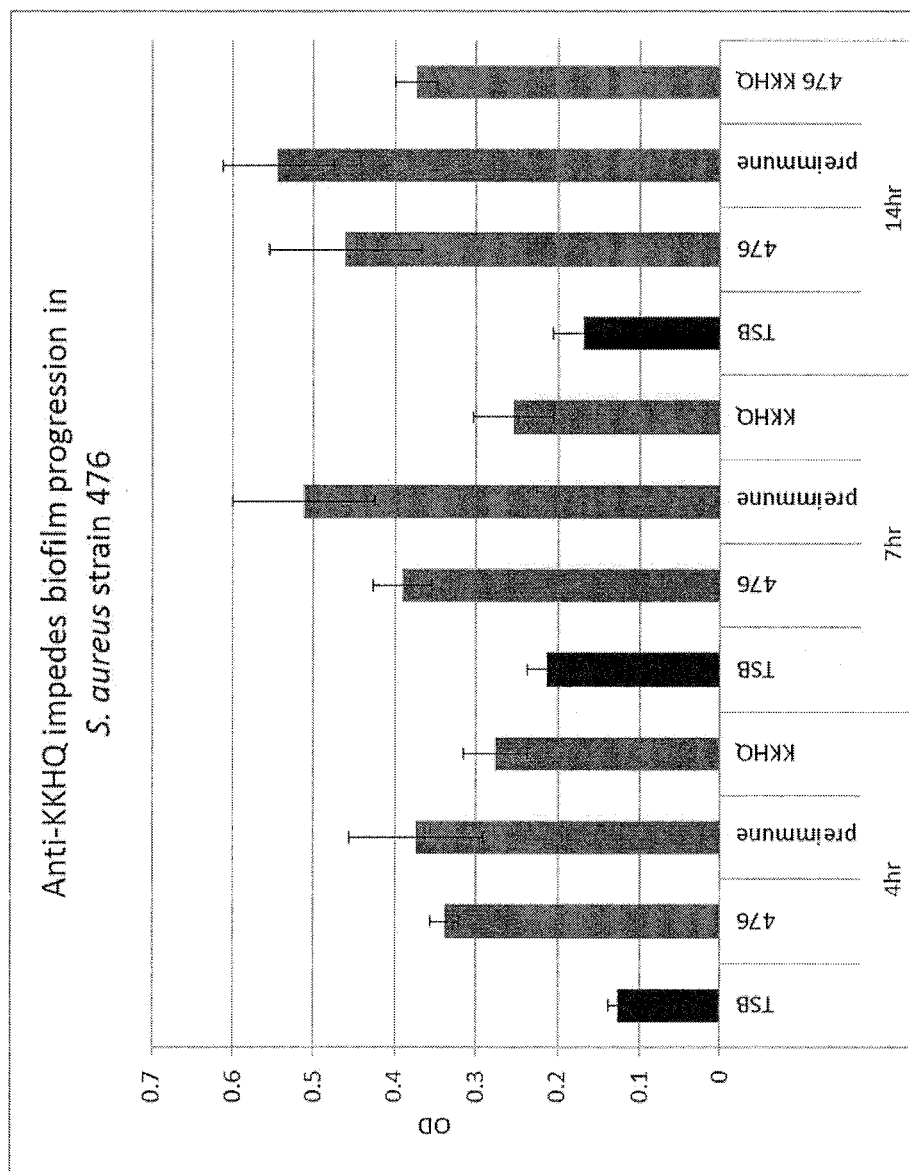
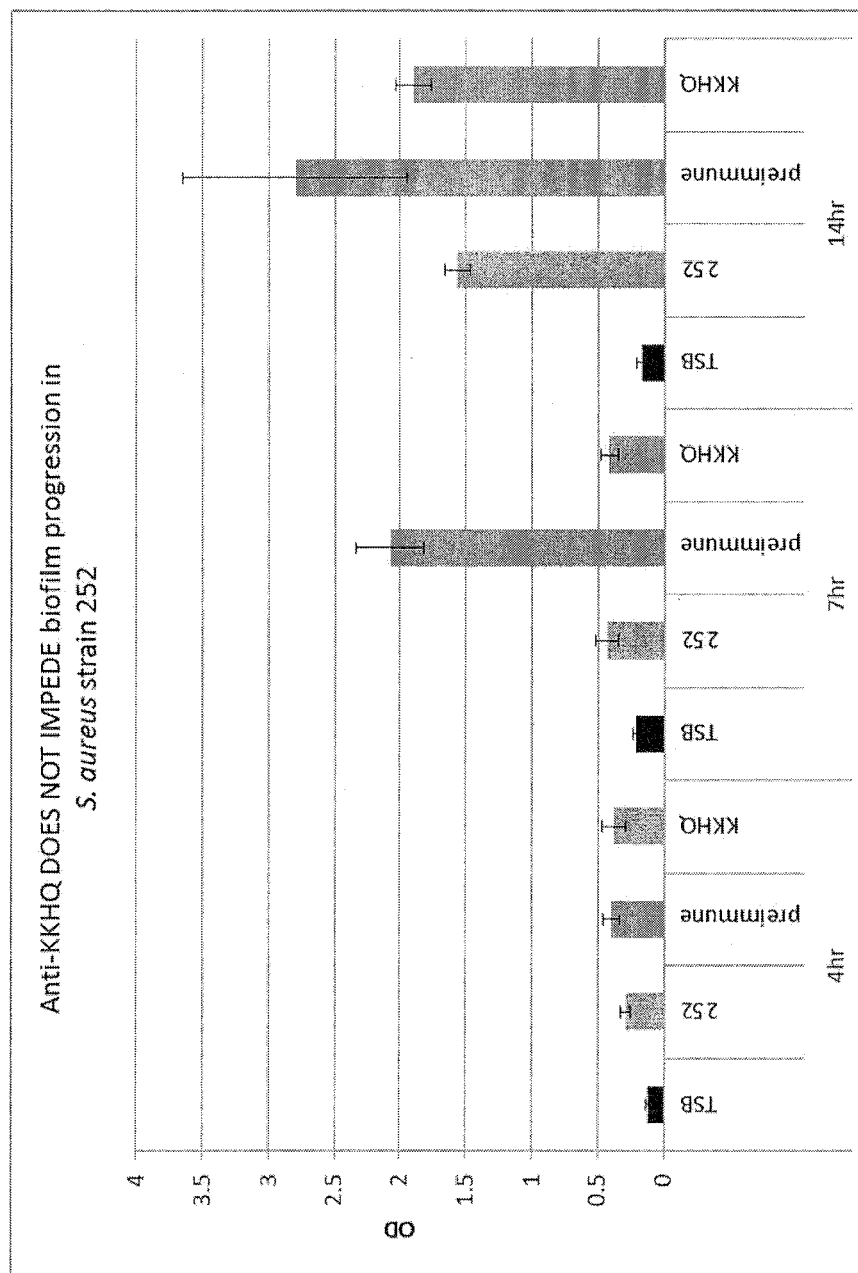


FIG. 12C



1

ANTIBODY BINDING MICROBIAL HEPARIN BINDING MOTIF TO RETARD OR PREVENT MICROBIAL BIOFILM FORMATION ON IMPLANTED MEDICAL DEVICES

This application is a Continuation-In-Part of co-pending PCT/US2013/031499 filed Mar. 14, 2013, which claims priority to U.S. application Ser. No. 61/636,243 filed Apr. 20, 2012, each of which is expressly incorporated by reference herein in its entirety.

A novel interaction between *Candida albicans* (*C. albicans*) surface proteins and heparin, a drug commonly used in patients at risk for catheter-associated biofilms and candidemia, is disclosed with an antibody that reduced or prevented a microorganism surface protein from binding to heparin, and reduced catheter-associated biofilm formation, where the reduction in biofilm formation can reduce the microorganism's virulence in a patient, with reduction encompassing any level of decrease.

Heparin is a highly sulfated, non-branched, anionic disaccharide composed of uronic acid, predominantly iduronic acid, in 1,4 linkage with glucosamine. Heparin is a known anticoagulant that is often infused in indwelling catheters to prevent the blood flowing through the catheter from clotting. Heparan sulfate is composed of the same disaccharide but is less sulfated and has a more varied structure than heparin. Heparan sulfate linked to core proteins, such as syndecans or glypicans, forms heparan sulfate proteoglycans (HSPGs). HSPGs are widely expressed on mammalian cell surfaces and in the extracellular matrix. Heparin binding motifs can bind to both heparin and heparan sulfate.

Several microbial proteins bind heparin and heparan sulfate, e.g., HIV-1 gp120, CypA, and Tat; hepatitis C envelope glycoprotein E2; herpes simplex virus glycoproteins B, C, and D; dengue virus envelope protein; *Plasmodium falciparum* circumsporozoite (CS) protein, *Listeria monocytogenes* ActA, and LcI of *Legionella pneumophila* (1-11). Such microbial protein interactions with surface HSPGs typically promote binding and entry of the microbial organism into the mammalian cell (except for HIV-1 gp120 and Tat), but the mechanisms by which individual microbial proteins recognize heparin or heparan sulfates are largely undefined. Positively charged tripeptides containing Lys or Arg in the amino terminus of *L. monocytogenes* ActA (aa 40-230) and in *L. pneumophila* LcI (aa 69-349) were shared with tripeptides in a number of heparin-binding mammalian proteins such as EGF-like growth factor and vonWillebrand factor, but the precise amino acids that mediated binding to heparin were not defined for ActA or LcI (10-11).

Eukaryotic proteins that bind heparin express either linear or conformational heparin binding motifs (HBM). Linear heparin binding motifs are short, conserved peptides containing both basic (B) and hydrophobic (X) amino acids in specific patterns first identified by Cardin (XBBXXBX) and Weintraub (XBBBXXBX) and subsequently expanded by Sobel (XBBBXXBBBXXBBX) (12-14). Molecular modeling studies suggested that basic amino acids such as lysine, arginine, and histidine were critical for interaction with anionic sulfate or carboxylate groups in heparin through electrostatic and hydrogen bonds. For example, substituting alanine for basic amino acids in linear HBM in the morphogen sonic hedgehog abolished binding to heparin (15). However, because many heparin-binding proteins failed to exhibit linear motifs, the concept of spatial orientation of basic residues was propounded (16). The CPC clip motif, a structural signature in which two cationic residues surround one polar residue, was conserved in eukaryotic heparin binding pro-

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teins, as well as in vaccinia complement protein and papillomavirus 18 capsid protein (17)

Because of its propensity to form biofilms in implanted medical devices such as indwelling central venous catheters, *Candida albicans* (*C. albicans*) is a leading pathogen in infections of indwelling catheters. A biofilm is a multilayered structure of microbes embedded in a polysaccharide matrix that forms on central venous catheters as well as implanted prosthetic joints, contact lenses, and other such medical devices (18, 19). A biofilm is undesirable because antibiotics and host defenses cannot penetrate it, thus a biofilm prevents elimination of microorganisms that have attached or adhered to a surface of an implanted medical device. For example, because of the high flow-through in a central venous catheter, sessile projections from a biofilm break off and are carried into the bloodstream to cause infection that can be carried by the blood to other sites in the body.

Heparin is a known anticoagulant, and catheters are often infused with heparin to prevent blood flowing through the catheter from clotting (20). Because heparin inhibited attachment of *C. albicans* to extracellular matrix proteins (21), the inventors hypothesized that *C. albicans* interacted with heparin through linear heparin binding motifs (HBMs): conserved sequences of basic (Arg, Lys, H is) and hydrophobic amino acids in prescribed patterns.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 schematically illustrates the internal lumen of an indwelling catheter.

FIGS. 2A-E demonstrate lack of an effect by heparin on wild type *Candida albicans* (*C. albicans*) planktonic growth and morphology.

FIGS. 3A-D demonstrate *C. albicans* binding to heparin and heparin analogs in vitro.

FIGS. 4A-C show that two lysine residues in linear heparin binding Motif 1 in *C. albicans* Int1 mediate binding to heparin, as measured by an ELISA assay in vitro. FIG. 4A discloses SEQ ID NOS 1, 40, 48, 2, 3, 52 and 53, respectively, in order of appearance. FIG. 4B discloses SEQ ID NO: 2. FIG. 4C discloses SEQ ID NOS 52-53, respectively, in order of appearance.

FIG. 5 compares pre-immune rabbit IgG with affinity-purified IgG raised against the sequence HKQEKQKKHQIHKV (SEQ ID NO:4) for ability to inhibit binding of *C. albicans* to heparin.

FIGS. 6A-C show consequences of heparin binding in vitro. FIG. 6A discloses SEQ ID NO: 65. FIG. 6B discloses SEQ ID NO: 66. FIG. 6C discloses SEQ ID NO: 67.

FIGS. 7A-B show consequences of heparin binding in vitro.

FIGS. 8A-F demonstrate that heparin binding motifs contribute to biofilm formation in vivo.

FIG. 9 demonstrates that affinity-purified IgG raised against the sequence HKQEKQKKHQIHKV (SEQ ID NO:4) substantially inhibited biofilm production in jugular venous catheters in rats. FIG. 9 discloses SEQ ID NO: 6.

FIG. 10 schematically illustrates the internal lumen of an indwelling catheter in the presence of antibody against heparin binding motifs on cell wall proteins of *C. albicans*.

FIG. 11 shows Anti-KKHQ specific binding to *Candida* species.

FIGS. 12A-C show Anti-KKHQ effect on biofilm progression.

Thirty-four *C. albicans* surface proteins encoding at least one linear heparin binding motif (HBM) were identified from a sequence-based search. Twelve of these 34 proteins flanked

recently described transcriptionally active regions that are involved in biofilm production. Thirty-three of these 34 proteins are known targets of master biofilm regulators (22).

C. albicans binding to 2.5 units (12.5 µg) solid-phase heparin in an ELISA assay was significantly decreased by desulfation of heparin at the 2-O or 2-N groups, and by preincubation of *C. albicans* with heparin. The protein Int1 contained the largest number of HBMs; Int1 is a surface protein that is involved in adhesion, filamentation, and antigenic recognition. In the Int1 sequence ₈₀₄QKKHQIHK (SEQ ID NO: 1), alanine mutation of lysine residues at positions 805/806 significantly reduced binding of the Int1 mutant to heparin. Rabbit IgG recognizing the polypeptide ₇₉₉HKQEKQKKHQIHKV₈₁₂ (SEQ ID NO: 4) inhibited *C. albicans* binding to heparin by 19%; pre-immune rabbit IgG had no effect.

Consequences of heparin binding in vitro included removal of *C. albicans* surface antigens such as Eno1, Pgk1, Tdh3, and Ssa1/2, which themselves contain putative HBMs; impairment of histatin-mediated killing; and modulation of gene expression. In vivo, substitution of alanine residues for lysines at positions 805/806 in ₈₀₄QKKHQIHK (SEQ ID NO: 1) markedly attenuated biofilm formation in central venous catheters in rats. In addition, pre-incubation of *C. albicans* with rabbit IgG recognizing the polypeptide ₇₉₉HKQEKQKKHQIHKV₈₁₂ (SEQ ID NO: 4) inhibited biofilm formation in vivo; pre-immune IgG had no inhibitory effect. These results identify linear HBM in *C. albicans* surface proteins, characterize specific lysine residues that mediate heparin binding, and demonstrate relevance for innate and adaptive immunity in vitro and biofilm formation in vivo.

The inventive method and composition ameliorated this undesirable situation by using antibodies against a specific heparin-binding motif that is expressed by surface proteins on *C. albicans*; other microorganisms also express similar heparin binding motifs. Without being held to a single theory, the disclosed antibodies reduce or prevent the undesirable biofilm from forming on the surface of a medical device, particularly a plastic device, implanted in a patient. Microorganisms include bacteria, yeast, fungi, etc. as known to one skilled in the art. Medical devices include those implanted or implantable in a patient, particularly plastic medical devices, including but not limited to catheters and central lines but excluding current non-plastic implanted joints, pacemakers, pacemaker wires, and spinal rods.

In one embodiment of the inventive method, antibodies were produced against a linear heparin binding motif that was expressed by a surface protein of *Candida albicans* (*C. albicans*). Related heparin binding motifs occur in surface proteins from *Staphylococcus epidermidis* (*S. epidermidis*) and *Staphylococcus aureus* (*S. aureus*).

Linear heparin binding motifs are sequential amino acids that conform to one of three consensus motifs (Cardin, Sobel, or Weintraub motifs); they have been identified in multiple mammalian proteins that bind heparin. While conformational heparin binding motifs also occur in some proteins, they must be identified by crystal structure of the protein. Antibodies to these linear heparin binding motifs inhibited *C. albicans* from adhering to heparin bound to a plastic surface (a microtiter plate).

Of the six million patients who have a central line implanted each year, more than 75,000 will develop bloodstream infections (23). Central line-associated blood stream infections (CLABSI) are a major source of hospital-acquired infection; over 43,000 CLABSI occurred among patients hospitalized in U.S. intensive care units (ICUs) in 2001, which is 3.2 infections for every 1000 line days. With a protocol for their sterile insertion and daily maintenance, CLABSI in

ICUs dropped to 18,000 in 2009, but even with these improvements in ICUs procedures, more than 23,000 CLABSI occurred in patients on inpatient wards and 37,000 CLABSI occurred in outpatient hemodialysis patients in 2009. Even with a 50% reduction in CLABSI in ICU patients, the annual cost of these infections across the U.S. is more than \$13 billion (23).

Staphylococcus epidermidis is the most common cause of CLABSI in central lines, with gram-negative rods second, *S. aureus* third; and *Candida* species fourth (24). In peripherally inserted central catheters (PICC), i.e., central lines inserted through peripheral veins in the arm, *Candida* species are the second most common cause of line infection (25). *Candida* spp. are equivalent to *S. aureus* in infections in central lines in the ambulatory setting (26). Among *Candida* spp., *C. albicans* is the most common cause of CLABSI (24).

When *Candida* species infect a central venous catheter, the organism enters the bloodstream to cause candidemia. Hosts at highest risk of candidemia, include burn patients and patients on a pump during coronary artery bypass procedures (27). Neutropenic oncology patients, premature newborns, and patients with major abdominal surgery such as intestinal resections are also high-risk patients with 8- to 16-fold the number of infections of other patient groups (27).

A major risk factor in candidemia is the presence of central lines (28-33), whose lumens are a site for formation of microbial biofilms that are effectively shielded from antimicrobial agents and host defenses. A second commonality is the use of heparin as an anticoagulant in most catheters. The amount of heparin is often considerable; e.g., a premature newborn in a neonatal intensive care unit may receive 150 units/kg/day, compared to 2.8 units/kg for a 70 kg adult receiving a 100 unit flush twice a day (20).

The ability to bind heparin is a unifying feature among *S. epidermidis*, *S. aureus*, and, *Candida albicans* (34-36), three of the leading causes of catheter-associated bloodstream infections. In experiments with *S. aureus*, high concentrations of heparin have been shown to increase *S. aureus* biofilm formation in vitro (37).

A first step in biofilm formation is microorganism adhesion or adherence to a surface of the implanted device, such as a catheter (38). In FIG. 1, negatively charged heparin molecules (circles) injected into the catheter lumen bind to the positively charged catheter surface, and heparin sulfate moieties (curved lines) are exposed on vascular endothelium lining catheter lumens. Linear heparin binding motifs present on the surface of microorganisms, such as *C. albicans*, interact either with heparin (A) or with heparin sulfate (B) and enable the organism to attach to the inside of the catheter. As the attached microorganisms replicate, they form the multi-layered structure of a biofilm (C) and secrete the polysaccharide mortar-like matrix that holds the biofilm together. The inventors hypothesize that interactions between heparin and surface proteins on microorganisms such as *C. albicans* facilitate biofilm formation, and subsequent infection of the bloodstream when projections from biofilm break off into the catheter lumen and enter the bloodstream.

Heparin is associated with central line infections, particularly those due to *Candida* spp. In a randomized trial of 260 central lines, 128 were coated with chlorhexidine and 132 were coated with heparin; all *Candida* colonization and all candidemias occurred in patients whose central lines were coated with heparin (39). A second study in renal dialysis patients showed that those patients who received 5000 units of heparin in the middle of dialysis had considerably more catheter loss due to infection compared to patients not treated

with heparin. Although staphylococcal infections were most common in this patient group, candidemia also occurred (40).

One mechanism by which heparin interacts with proteins is through linear heparin binding motifs, conserved sequences of basic amino acids such as lysine, arginine, and histidine interspersed with hydrophobic amino acids (14). Microbial heparin-binding proteins are identified by short consensus motifs of basic (B) and hydrophobic (X) amino acids, as defined by Cardin [XBBXB], Weintraub [XBBBXXB], or Sobel [XBBBXXBBBXXB].

In the invention, a computer-based algorithm examined all 400 *C. albicans* surface and cell wall proteins for heparin binding motifs (HBM). Of the 6,000 known *Candida albicans* proteins, about 400 are localized to the cell wall of yeast or hyphae, depending upon the extraction technique (41). A sequence-based search identified putative Cardin, Sobel, or Weintraub motifs in 159 *C. albicans* proteins. Table 1 shows HBM in cell surface proteins expressed by *C. albicans*. The 34 proteins in Table 1 include only those that have the attribution "cell surface", "fungal cell wall", "yeast cell wall", or "hyphal cell wall" as the cellular component in the *Candida* Genome Database, where * = Weintraub motif; # = Cardin motif; ‡ = Identified by molecular modeling studies; A = "cell wall" in protein description (as of January 2011); obtained from *Candida* Genome Database (CGD); B = "cell wall" in Gene Ontology (GO) annotation (as of January 2011); obtained from CGD; C = cell wall proteins as reviewed by Alberti-Segui (42) and D = cell wall proteins as reviewed by Chaffin (41).

The following procedure was used to identify a set of cell wall proteins in *C. albicans* with HBM. The first step combined evidence to identify possible cell wall proteins in *C. albicans*. Specifically, using "cell wall" as keywords, 154 putative cell wall proteins were identified from protein descriptions, and another 245 unique cell wall proteins were identified from Gene Ontology functional annotations from *Candida* Genome Database (www.candidagenome.org; Assembly 21). Also included were 125 non-redundant cell wall proteins identified by Alberti-Segui (50) and another 174 reported by Chaffin (22) and mapped annotations to Assembly 21. The second step screened for consensus HBM in protein ORF sequences of these cell wall proteins from Assembly 21. Three types of consensus motifs were included: Weintraub (XBBBXXB), Sobel (XBBBXXBBBXXB), and Cardin (XBBXB) motifs, where B is a basic amino acid and X is a hydrophobic amino acid. In the algorithm, basic amino acids are H, K, and R. Hydrophobic amino acids are W, F, Y, L, I, C, M, G, V, S, T, A, N, P, and Q. A total of 159 cell

wall proteins were identified that have possible HBM, and some have multiple motifs. The 159 proteins that met the search criteria were then manually curated to select only those proteins whose cell wall localization was confirmed by manual or computational methods in the *Candida* Genome Database.

Proteins with GPI anchoring sequences within these 159 cell wall proteins with HBM were further identified. Three Web servers were used to perform GPI anchoring sequence prediction: SignalP (43), GPI-SOM (44), and PredGPI (45). Combined results of these three prediction methods indicated that 15 out of these 159 proteins had GPI anchoring sequences, four of which overlapped with reviewed GPI anchoring proteins (46).

The algorithm identified 34 cell wall proteins with potential linear heparin binding sequences that matched the Cardin, Weintraub, and Sobel motifs. The 34 identified cell wall proteins were Ahp1, Als7, Atcl, Bud2, Cat1, Cef3, Chs1, Crh12, Dot4, Not5, Pdi1, Pga4, Pgl1, Phr3, Rbt1, Rps6A, Sam2, Srb1, Ssa2, Ssb1, Tdh3, Eft2, Eno1, Gap1, Gph1, Gpm2, Hem13, Hsp104, Hsp70 (also called Ssa1), Ino1, Int1, Ipp1, Ugp1, Xyl2. Three of these proteins, Als7, Pga4, and Rbt1, have GPI anchors. The genes encoding eleven of these proteins, Als7, Cat1, Dot4, Eno1, Gph1, Ino1, Rbt1, Sam2, Srb1, Ssa2, and Ugp1, are located in newly defined transcriptionally active regions that are critically involved in biofilm formation (22). Moreover, many of these proteins are known to be regulated by at least one of six master biofilm regulators; by RNA-seq, where RNA-seq is defined as the use of high-throughput sequencing techniques to sequence cDNA in order to get sequence information about the transcriptome, the sample's RNA content; CAT1, GAP1, GPH1, and HSP104 are up-regulated in biofilm formation, while CHS1, ENO1, GPM2, HEM13, INO1, IPP1, PGK1, RPS6A, SAM2, SRB1, SSA2, SSB1, and TDH3 are down-regulated (22).

Given the presence of putative HBM among *C. albicans* surface proteins and the possible interactions with heparin in central venous catheters or with heparin sulfate proteoglycans expressed on host tissues such as vascular endothelium, the biochemical determinants and immunologic consequences of this interaction were defined. The protein Int1 (SEQ ID NO: 5) (accession number P53705.2; GI 187608862) had the highest number of HBM (five). Int1 (SEQ ID NO: 5) spans 1711 amino acids and is localized to the cell wall of the bud neck in *C. albicans*. It mediates adhesion, hyphal formation, and virulence, defined as the ability to cause disease such as bloodstream infection (47). Using the same search technique, HBM were also detected in surface proteins of *S. epidermidis* and *S. aureus*.

TABLE 1

Gene ID (Assembly 21)	Protein	Ref.	# of motifs	Motif 1	Motif 2	Motif 3	Motif 4	Motif 5
orf19.4257	Int1	C	5	804QKKHQIHK [‡] (SEQ ID NO: 1)	1383THKGRF [#] (SEQ ID NO: 40)	1530MKRGKP [#] (SEQ ID NO: 48)	1593FKKRFKKL* (SEQ ID NO: 2)	1612SHKTRA [#] (SEQ ID NO: 3)
orf19.1738	Ugp1	B, D	3	176SHRIRV [#] (SEQ ID NO: 7)	310IKKFKY [#] (SEQ ID NO: 41)	368IRHFKG [#] (SEQ ID NO: 49)		
orf19.3370	Dot4	B	3	442NKKGKS [#] (SEQ ID NO: 8)	519CHKCHN [#] (SEQ ID NO: 42)	635PKRFPK [#] (SEQ ID NO: 50)		
orf19.3651	Pgk1	A, B, D	3	136GKKVKA [#] (SEQ ID NO: 9)	146VKKFRQ [#] (SEQ ID NO: 43)	168AHRAHS [#] (SEQ ID NO: 51)		
orf19.4660	Rps6A	B, D	2	185QRKRALK* (SEQ ID NO: 10)	192AKKVKN [#] (SEQ ID NO: 44)			

TABLE 1 -continued

Gene ID (Assembly 21)	Protein	Ref.	# of motifs	Motif 1	Motif 2	Motif 3	Motif 4	Motif 5
orf19.5107	Not5	B	2	¹¹⁴ QKRSRF [#] (SEQ ID NO: 11)	³³³ VKKLKP [#] (SEQ ID NO: 45)			
orf19.5130	Pdi1	D	2	²¹⁰ NKKFKN [#] (SEQ ID NO: 12)	³⁰¹ GKKYRG [#] (SEQ ID NO: 46)			
orf19.6387	Hsp104	D	2	⁵³ VKRARY [#] (SEQ ID NO: 13)	¹⁹⁹ ARRSKS [#] (SEQ ID NO: 47)			
orf19.1065	Ssa2	B, D	1	²⁵⁸ LRRLRT [#] (SEQ ID NO: 14)				
orf19.1067	Gpm2	B	1	⁴⁵ IKKNHL [#] (SEQ ID NO: 15)				
orf19.1327	Rbt1	A, B, C	1	¹²¹ GKKVKQ [#] (SEQ ID NO: 16)				
orf19.2762	Ahp1	B, D	1	¹⁷¹ LKRIHN [#] (SEQ ID NO: 17)				
orf19.2803	Hem13	B, D	1	²⁵⁷ IRRGY [#] (SEQ ID NO: 18)				
orf19.3590	Ipp1	B, D	1	⁷⁴ TKKGKL [#] (SEQ ID NO: 19)				
orf19.377	Phr3	B, C	1	¹¹⁵ PHHHLNRY [*] (SEQ ID NO: 20)				
orf19.395	Eno1	B, D	1	¹⁴¹ AKKGKF [#] (SEQ ID NO: 21)				
orf19.3966	Crh12	A, B, C	1	⁴²⁰ TKHIHN [#] (SEQ ID NO: 22)				
orf19.4035	Pga4	B, C, D	1	²⁵⁹ AKRPRP [#] (SEQ ID NO: 23)				
orf19.4152	Cef3	B	1	⁶²³ LRKYKG [#] (SEQ ID NO: 24)				
orf19.4304	Gap1	B	1	⁷² QRKLKT [#] (SEQ ID NO: 25)				
orf19.4980	Ssa1	B, D	1	²⁵⁹ LRRLRT [#] (SEQ ID NO: 26)				
orf19.5188	Chs1	B, C	1	¹²⁶ PKRQKT [#] (SEQ ID NO: 27)				
orf19.5788	Eft2	B, D	1	⁵⁸¹ NKHNRI [#] (SEQ ID NO: 28)				
orf19.6190	Srb1	B, D	1	¹²³ FHKAHG [#] (SEQ ID NO: 29)				
orf19.6214	Atc1	A, B	1	⁹⁵⁹ PKRVKV [#] (SEQ ID NO: 30)				
orf19.6229	Cat1	B	1	⁸² GKKTRI [#] (SEQ ID NO: 31)				
orf19.6367	Ssb1	B, D	1	²⁶³ LRRLRT [#] (SEQ ID NO: 32)				
orf19.657	Sam2	B, D	1	³⁸⁰ PKKLKF [#] (SEQ ID NO: 33)				
orf19.6814	Tdh3	B, D	1	⁷⁰ GHKIKV [#] (SEQ ID NO: 34)				
orf19.7021	Gph1	B, D	1	⁶⁵⁶ TKHHIPKA [*] (SEQ ID NO: 35)				

TABLE 1 -continued

Gene ID (Assembly 21)	Protein	Ref.	# of motifs	Motif 1	Motif 2	Motif 3	Motif 4	Motif 5
orf19.7400	Als7	B, C	1	¹⁴⁸² SKRNKN [#] (SEQ ID NO: 36)				
orf19.7585	Ino1	B, D	1	¹⁶¹ MKRAKV [#] (SEQ ID NO: 37)				
orf19.7676	Xyl2	A, B, D	1	³³⁰ THRFFK [#] (SEQ ID NO: 38)				
orf19.940	Bud2	B	1	⁷¹⁶ LRKGKS [#] (SEQ ID NO: 39)				

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Heparin effect on *Candida albicans* growth and morphology was determined. A single colony of BWP17 wt was inoculated into 3 ml YPD medium and incubated overnight at 30° C. with shaking at 225 rpm. Overnight cultures were diluted to an OD₆₀₀ of 0.1 in 5 ml of YPD (yeast) or RPMI-HEPES (hyphae) in a 50 ml conical polypropylene tube and incubated in the presence or absence of 100 units/ml preservative-free pharmaceutical heparin at 30° C. for six hours. Cells (1 ml) were fixed with an equal volume of 4% formaldehyde in FACS buffer at 4° C. for one hour, then pelleted (7,000 rpm for 3 minutes) and washed with PBS. After reconstitution in 0.5 ml PBS, filipin (50 mg/ml stock solution in DMSO) was added to a final concentration of 100 µg/ml and incubated with cells at room temperature for five minutes. Cells were pelleted, washed with PBS and mounted on a slide using Fluoromount G. For calcofluor white staining, after six hours incubation, 1 µl of 1 mg/ml stock solution of calcofluor white in 0.1 N NaOH was added to 100 µl cells to a final calcofluor white concentration of 10 µg/ml.

Microscopy was performed by imaging filipin, PKH26, DAPI, and heparin-Alexa Fluor 488 on a Nikon Ti-E inverted microscope with a 100×CFI APO oil NA 1.49 objective. Filipin was excited with a Prior Lumen 200 metal halide light source set at 10% light output. This light was further attenuated by ND4 and ND8 neutral density filters in series to reduce light output to approximately 3% of output from the liquid light guide. The filters used for imaging were EX 360/40, dichroic 400 nm Ip, EM 460/50. Images were acquired with an Andor iXon emccd camera. Exposure times were 391 ms. Excitation of PKH26/DAPI/Heparin-Alexa Fluor 488 triple staining was accomplished with a Nikon A1R si laser scanning confocal. DAPI was imaged with 405 nm excitation and a 450/50 filter with a laser power of 16.3 and a photomultiplier tube (pmt) voltage of 92. Alexa Fluor 488 was imaged with 488 nm excitation from an Argon-ion laser and a 525/50 filter with a laser power of 6.0 and a pmt voltage of 84. PKH26 was imaged with 561 nm excitation and a 595/50 filter with a laser power of 13.2 and a pmt voltage of 108. Gains were kept below 110 to eliminate contribution from *C. albicans* autofluorescence; lack of autofluorescence was confirmed by comparison to a control sample without heparin-Alexa Fluor 488. Images were processed with Nikon NIS-Elements AR 4.11.00 64-bit software. Calcofluor white slides were examined using a Zeiss Axiovert 200M fluorescent microscope equipped with a Plan-Apochromat 63×/1.40 oil objective lens with 1.6× optivar and DAPI filter at 350 nm excitation and 460 nm emission. Images were captured using a Zeiss AxioCam color camera and processed with Axiovision version 4.8.2.

Flow cytometry was performed by inoculating a single colony of BWP17 wt into 3 ml YPD medium and incubating

overnight at 30° C. with shaking at 225 rpm. Overnight cultures were diluted to OD₆₀₀ of 0.1 in 5 ml of YPD or RPMI-HEPES and incubated in the presence or absence of 100 units/ml preservative-free pharmaceutical heparin at 30° C. for one hour. One ml aliquots of each mixture were removed, pelleted (7,000 rpm for 3 min), washed twice with PBS, and reconstituted in 1 ml PBS. Flow cytometry analysis was performed using the Imagestream^X flow cytometer (Amnis, Seattle Wash.) equipped with a 405 nm, 488 nm, 653 nm, laser and multi-mag function. The 40× magnification, 10 mm/sec flow rate, and 488 nm laser were used to collect SSC and Brightfield parameters. The flow cell allows particles up to 100 µm wide (height unlimited) to be collected in the instrument. INSPIRE (v.6.0) software was used to acquire events. Software analysis using IDEAS (v5.0) identified percent hyphae using aspect ratio, height and width features using the side scatter (Channel 6) and brightfield (Channel 1) parameters. Objects were selected within each file and tagged to use as identification of truth sets within the population.

For growth curves in the presence or absence of soluble heparin, a single colony of BWP17 wt was inoculated into 3 ml YPD medium and incubated overnight at 30° C. with shaking at 225 rpm. Overnight cultures were diluted to an OD₆₀₀ of 0.1 in 5 ml of YPD, RPMI-MOPS, RPMI-HEPES or CSM in a 50 ml conical polypropylene tube. Preservative-free pharmaceutical heparin (1000 units/ml) was added to yield a final concentration of 100 units/ml in the heparin-treated samples. Cultures were grown at 30° C. with shaking and OD₆₀₀ measured periodically. Doubling times (t_d) were calculated based on the equation $t_d = \ln 2 / \mu$ (μ =specific growth rate (48)). One hundred units/ml heparin, the concentration recommended to prevent clotting of central venous catheters (20), had no effect on doubling times of planktonic yeast cells grown in YPD, RPMI-HEPES, RPMI-MOPS or CSM, as shown in FIGS. 2A and 2B.

FIGS. 2A-E demonstrate lack of an effect by heparin on wild type *Candida albicans* (*C. albicans*) planktonic growth and morphology.

FIG. 2A shows that growth curves in the presence (open circles) or absence (closed diamonds) of soluble heparin were identical.

FIG. 2B shows that doubling times in various media at 30° C. in the absence and presence of heparin (100 units/ml) were identical.

FIG. 2C shows that percent hyphae as determined by flow cytometry did not differ after incubation with or without heparin (100 units/ml) under conditions that favor yeast (YPD, 30° C.) or hyphae (RPMI-HEPES, 37° C.).

FIGS. 2D-E show that the integrity of membrane sterols identified with filipin (FIG. 2D) and the location of septin rings identified with calcofluor white staining (FIG. 2E) were

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identical between untreated and heparin-treated organisms. Thus, concentrations of heparin commonly used to maintain patency of central venous catheters (100 units/ml) did not lead to readily observable differences in growth or cellular architecture of the organism.

Binding of heparin by *C. albicans* was examined in vitro. Heparin was labeled with Alexa Fluor 488 by a method modified from Osmond (49). Briefly, 5 mg heparin (Sigma) was dissolved in 0.5 ml MES buffer (0.1M 2-(N-morpholino) ethanesulfonic acid hydrate, pH 4.5, Sigma), then mixed with a solution of 1 mg Alexa Fluor 488 hydrazide (Life Technologies) in 0.4 ml MES buffer. After adding 0.2 ml of EDC solution (15 mg 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide/ml water, Thermo Scientific), the mixture was stirred at room temperature for 30 min. A second 0.2 ml aliquot of EDC solution was added and the mixture stirred at room temperature for an additional 30 min. After adding 1.3 ml NaOAc (1 M, pH 4.8) and stirring at room temperature for one hour, heparin-Alexa Fluor 488 was purified on a PD-10 desalting column (GE Healthcare) equilibrated in autoclaved nanopure water. Fractions containing the labeled material detected at 490 nm were combined and dried overnight on a SpeedVac concentrator (Savant) with heating. The resulting solid was dissolved in autoclaved nanopure water to 10 mg/ml and stored at 4° C.

Heparin-Alexa Fluor 488 bound to PKH26- and DAPI-labeled *C. albicans*. A single colony of BWP17 wt was inoculated into 3 ml YPD medium and incubated overnight at 30° C. with shaking at 225 rpm. Using a PKH26 Red Fluorescent Cell staining kit (including PKH26 dye and diluent C, Sigma), 2×10^7 cells were mixed with 1 ml diluent C. To this solution was added a mixture of 4 μ l PKH26 dye in 1 ml diluent C, and the mixture allowed to stand at room temperature for 3 min. After adding 10 ml 3% BSA, the mixture was centrifuged at 3,000 rpm for 7 min. The resultant magenta-colored cells were washed with 10 ml each of 1% BSA and PBS, with centrifugation after each step. After supernatant removal, the cells were reconstituted in 10 ml fresh PBS. Two 1 ml (2×10^6 cells) aliquots were placed in separate 15 ml conical tubes, centrifuged, and supernatants removed. Each was reconstituted with 2 ml YPD, and heparin-Alexa Fluor 488 (0.1 ml of above 10 mg/ml solution) was added to the experimental sample. Control and experimental samples were incubated at 30° C. with shaking (225 rpm) for 30 min, at which time 0.4 ml aliquots of each were removed, pelleted (10,000 rpm for 3 min), and washed twice with PBS. After reconstitution with 0.5 ml PBS, DAPI (4',6-diamidino-2-phenylindole dihydrochloride (Sigma, stock solution of 5 mg/ml) was added to a final concentration of 1 μ g/ml and the solution allowed to stand at room temperature for 10 min. Cells were pelleted and washed twice with PBS, then mounted on a microscope slide using Fluoromount G.

To measure the binding of *C. albicans* to solid-phase heparin, a heparin-binding ELISA assay was developed, as follows. A single colony of each *C. albicans* wild type or mutant strain from a YPD plate was inoculated into 3 ml YPD medium and incubated overnight at 30° C. with shaking at 225 rpm. Sigma heparin (fresh solution made daily), diluted to 25 units/ml in autoclaved, sterile-filtered PBS was added as 0.1 ml aliquots to each well of an allyl amine-coated 96-well heparin binding microtiter plate (BD Biosciences). The plate was incubated at room temperature overnight in the dark (50). In the morning, the plate was washed with acetate buffer (100 mM NaCl, 50 μ M NaOAc, 0.2% Tween 20, pH 7.2), incubated with 3% bovine serum albumin (BSA) in PBS at 30° C. for 1 hour, then washed with PBS. In the meantime, overnight cultures of *C. albicans* were subcultured by dilution to OD₆₀₀

of 0.2 in 25 ml of YPD, and grown at 30° C. to mid-log phase (OD₆₀₀ 0.6-0.7). Cells were pelleted (3,000 rpm for 7 minutes), washed twice with PBS and reconstituted in RPMI-HEPES to 4×10^5 , 2×10^5 , and 1×10^5 CFU/ml, respectively. One hundred μ l of each *C. albicans* dilution was applied to the microtiter wells; experiments for each strain were performed in quadruplicate. The plate was incubated at 30° C. for one hour and washed with PBS to remove non-adherent *C. albicans*. One-tenth ml of a biotinylated polyclonal rabbit anti-*C. albicans* antibody raised against soluble proteins in a *C. albicans* lysate (Meridian Life Science), which had been diluted 1:2500 in FACS-Tween (0.05% Tween 20 in FACS buffer consisting of 0.3% BSA), was added to each well. The plate was incubated at 30° C. for one hour, then washed with PBS-Tween (0.05% Tween 20 in PBS). One-tenth ml streptavidin alkaline phosphatase (Biolegend) diluted 1:10,000 in FACS-Tween was then added to each well, and the plate was incubated at 30° C. for 30 minutes. After washing with PBS-Tween and AKP buffer (100 mM Tris base, 50 μ M MgCl₂, 100 mM NaCl, pH 9.5.), 0.1 ml alkaline phosphatase substrate (KPL) was added to each well. After 45 minutes, absorbance at 595 nm was read on a Beckman Coulter DTX 880. When desulfated heparin analogs, chondroitin sulfate, or dermatan sulfate were used, equimolar amounts of the analogs (with respect to heparin) were applied to the allyl amine-coated 96-well plate instead of heparin.

The ability of soluble heparin to inhibit *C. albicans* binding to plate-bound heparin was demonstrated by pre-incubating 1×10^6 CFSE-labeled *C. albicans* with 200 units heparin at 37° C. for one hour in wells of a 96-well black plate. *C. albicans* was labeled with carboxyfluorescein succinimidyl ester (CFSE), succinimidyl ester (CFSE; Invitrogen), which is used to track cell division and therefore does not kill the organism (28), as follows. A single colony of *C. albicans* was suspended in CSM to an OD₆₀₀ of 0.1, diluted 1:200 into 25 ml CSM and grown at 30° C. overnight to mid-log phase (OD₆₀₀ 0.6-0.7). Cells were washed twice in sterile PBS and suspended in PBS at 2×10^7 CFU/ml. Carboxyfluorescein succinimidyl ester (Life Technologies) was dissolved in DMSO, diluted in PBS to 50 μ M, and 0.5 ml CFSE mixed with 0.5 ml *C. albicans* suspension. The mixture was incubated for 20 min at room temperature on a rotator, washed once with FACS buffer, once with PBS, then suspended in 1 ml FACS buffer. Intensity of labeling was determined as the mean fluorescence intensity using a BD Accuri C6 cytometer (San Jose, Calif.) with excitation at 488 nm and a 533/30 emission filter. Wells of a black 96-well microtiter plate were incubated with 0.1 mg poly-D-lysine for one hour at room temperature, washed three times with PBS, then incubated with 200 units of heparin in 100 μ l RPMI-HEPES overnight at room temperature. The following afternoon, *C. albicans* wild type strain and mutants were labeled with CFSE as follows: after growth of *C. albicans* to mid-log phase in CSM medium, organisms were washed two times in PBS and suspended in PBS at a concentration of 2×10^7 /ml. 0.5 ml cells was mixed with 0.5 ml CFSE, covered in foil, and rotated on a mixer for 20 minutes at room temperature. CFSE-labeled *C. albicans* cells were pelleted in a minifuge at 3000 rpm for four minutes, washed once with FACS buffer (0.3% bovine serum albumin in PBS), washed again with PBS, and then suspended in 1 ml FACS buffer at a concentration of 1×10^7 /ml. 100 μ l CFSE-labeled *C. albicans* (1×10^6) were deposited in test wells for 60 minutes at room temperature. At the end of the incubation period, wells were washed three times with PBS, and fluorescence in each well was measured (Beckman Coulter Multimode detector DTX880).

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FIGS. 3A-D demonstrate *C. albicans* binding to heparin in vitro. FIG. 3A shows confocal microscopy of *C. albicans* in the presence of heparin-Alexa Fluor 488, with (i) *C. albicans* cell wall stained with PKH26, (ii) nucleus stained with DAPI, (iii) cell wall outlined by heparin-Alexa Fluor 488, and (iv) arrows indicating co-localization of heparin with *C. albicans* cell surface. Incubation of *C. albicans* with 100 units/ml heparin-Alexa Fluor 488 for 30 mins at 30° C. demonstrated co-localization of heparin with the *C. albicans* cellular surface (FIG. 3A). Heparin deposition was seen at the interface of adjoining yeast cells and in individual cells.

FIG. 3B shows binding of 10,000 colony-forming units (CFU) *C. albicans* (OD₅₉₅) to increasing concentrations of heparin immobilized on an allyl amine-coated 96-well micro-titer plate; values are \pm SD of duplicate wells. Binding of *C. albicans* at 10,000 CFU/well did not change over heparin concentrations ranging from 1.25 units/well (6.3 μ g/well) to 20 units/well (100 μ g/well) (FIG. 3B), indicating saturation of the allyl amine plate by heparin in low concentrations.

FIG. 3C shows the linear dose-response for the heparin binding ELISA assay with 2.5 units/well heparin and increasing *C. albicans* input from 10,000 (left-most bar), 20,000 (center bar), and 40,000 (right-most bar) CFUs; the graph represents mean \pm SD of four experiments, *p<0.007 for all inputs. FIG. 3C shows that the OD₅₉₅ of the ELISA assay increases linearly as input of *C. albicans* increased from 10,000 colony forming units (CFU)/well to 40,000 CFU/well.

FIG. 3D shows binding of *C. albicans* (40,000 CFU/well) to equimolar amounts of heparin analogs desulfated at the 2-O (second from the left bar), 2-N (second from the right bar), or 6-O (right-most bar) positions versus heparin control (left-most bar) (normalized to 100%), with the graph representing mean \pm SD of three experiments, performed in triplicate *p \leq 0.003, **p=0.20 vs. heparin control. With equimolar amounts of desulfated heparins, binding of *C. albicans* was decreased by 11% when heparin was desulfated at the 2-O position of iduronic acid (p=0.003), and by 21% when heparin was desulfated at the 2-N position of glucosamine (p=0.002) (FIG. 3D). Removal of the sulfate group at the 6-O position of glucosamine did not significantly reduce binding.

Pre-incubation of CFSE-labeled *C. albicans* with 100 units/ml heparin decreased binding to heparin by 27% (p=0.022; data not shown). *C. albicans* also recognizes the related structures of chondroitin sulfate and dermatan sulfate.

Binding of *C. albicans* to heparin and to chondroitin sulfate was equivalent, but binding to dermatan sulfate was reduced (p=0.019; data not shown). These results confirmed that the binding of *C. albicans* to heparin and similar glycosaminoglycans (chondroitin sulfate and dermatan sulfate) could be reproducibly measured and, in the case of heparin, specifically inhibited by pre-incubating the organisms with heparin.

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These results also suggested that surface components of *C. albicans* may mediate *C. albicans* binding to heparin.

Location of HBM in Int1 (SEQ ID NO: 5) were determined. Five overlapping polypeptides spanning amino acids 51-1711 of Int1 were expressed with a 6xHis tag (SEQ ID NO: 54) in *S. cerevisiae* BJ3501 and purified by affinity chromatography (HisTrap column; GELifesciences). Fractions containing the His-tagged polypeptide were pooled, diluted with loading buffer (10 mM phosphate, pH 7.0 plus 250 mM NaCl) and applied to a heparin sepharose column (HiTrap Heparin HP; GE Lifesciences). Heparin-binding polypeptides were eluted with a step gradient of NaCl (0.5-2 M) in loading buffer. Eluted fractions were analyzed by SDS-PAGE and immunoblot using anti-His tagged antibody (Santa Cruz) and chemiluminescent detection (SuperSignal West Pico Mouse IgG Detection Kit, Pierce) according to manufacturers' instructions. Polypeptides spanning aa 656-1193 and aa 1548-1711 bound to heparin-Sepharose and were eluted with NaCl, indicating that the HBM in those domains, schematized as Motif 1 (SEQ ID NO: 1), Motif 4 (SEQ ID NO: 2), and Motif 5 (SEQ ID NO: 3) (FIG. 4A), were candidates for mediating binding to heparin. Bolded letters represent basic amino acids in heparin binding motifs; unbolded letters represent hydrophobic residues. Polypeptides spanning aa 51-385, aa 385-659, and 1188-1551 failed to bind to a heparin-Sepharose column.

The linear polypeptide spanning aa 656-1193 contains one potential heparin binding site, ⁸⁰⁴QKKHQIHK (SEQ ID NO: 1) (basic residues shown in red) (Motif 1 in Table 1 and FIG. 4A). The linear polypeptide spanning aa 1548-1711 contains a canonical Weintraub motif ¹⁵⁹³FKKRFFKL (SEQ ID NO: 2) (Motif 4 in Table 1 and FIG. 4A) and a canonical Cardin motif ¹⁶¹²SHKTRA (SEQ ID NO: 3) (Motif 5 in Table 1 and FIG. 4A). Sequence homology search using BLAST indicated that aa 1548-1711 contains a Pleckstrin homology domain (PHD), which is structurally resolved. Using homology-based 3-D modeling, we found that the three lysine residues and single arginine residue in Motif 4 were located on the rim of the PHD and might facilitate binding to a strong anion such as heparin by electrostatic interaction (FIG. 4B, arrows). The Cardin motif (Motif 5) did not share this conformation.

To test whether ⁸⁰⁴QKKHQIHK (SEQ ID NO: 1) (Motif 1) and/or ¹⁵⁹³FKKRFFKL (SEQ ID NO: 2) (Motif 4) mediated the binding of heparin to *C. albicans*, standard PCR-mediated mutagenesis (51) was used to construct a set of isogenic INT1 disruptants and mutants (Table 2).

TABLE 2

Strain	Abbreviation	Genotype	Source
BWP17	—	ura3::imm434/ura3::imm434 his1::hisG/his1::hisG arg4::hisG/arg4::hisG	[27]
BWP17WT	WT	BWP17 plus arg4::ARG4::URA3/his1::hisG::HIS1	[27]
VBDM2	—	as BWP17 plus int1::ARG4/int1::URA3	[60]
VBDM6-2	DD	as VBDM2 plus his1::hisG/HIS1	[60]
KO503	Motif 4	VBDM2 plus his1::hisG::HIS1-INT1 (KK1595AA)	current
KO507	Motif 1	VBDM2 plus his1::hisG::HIS1-INT1(KK804AA)	current
KO508	Motifs 1&4	VBDM2 plus his1::hisG::HIS1-INT1(KK804AA, KK1595AA)	current
KO509	Reint	VBDM2 plus his1::hisG::HIS1-INT1 (WT)	current

Construction of mutants was as follows. *C. albicans* genomic DNA was isolated from saturated overnight cultures using glass beads as described (52). A lithium acetate method was used to transform *C. albicans* (41). Plasmids and PCR products were purified using kits (Fermentas/ThermoFisher, Pittsburgh Pa.) or established methods (53). Pfu enzyme (New England Biolabs) with High Fidelity buffer was employed for

all amplifications. Products were sequenced to affirm fidelity prior to use. Primers are described in Table 3.

units (12.5 μ g) heparin in 100 μ l phosphate buffered saline (PBS) per well. The plate was covered and allowed to stand at

TABLE 3

Primer	Internal reference Sequence	Purpose
1	SAC2035UP gggagctcGTTACTTGTCTAATTAAGTTACTTCC (SEQ ID NO: 55)	SacI 5' INT1
2	MLU3' UTR ggacgcgtTTTTATCTTTTATGTAAATATATACTA (SEQ ID NO: 56)	MluI 3' INT1
3	INT1 KR1595AA F ATTGTCCAATTTTAAAGGCTGCTTTTTCAAATTAAT GGG (SEQ ID NO: 57)	mutate Motif 1
4	INT1 KR1595AA R CCATTAATTTGAAAAAGCAGCCTTAAAAATTGGAC AATC (SEQ ID NO: 58)	mutate Motif 1
5	INT1 KK805AA F GCATAACAAGAAAGCAGGCCGCCATCAAATTC ATAAAGTTCC (SEQ ID NO: 59)	mutate Motif 4
6	INT1 KK805AA R GGAACTTTATGAATTTGATGGCGGCCTGCTTTTC TTGTTTATGC (SEQ ID NO: 60)	mutate Motif 4
7	GHISR CTCCCGGCCCATGGCCGC (SEQ ID NO: 61)	check integration
8	HIS3AMP GTTGGTGTGGCCAGAGACTCT (SEQ ID NO: 62)	check integration

A single copy of Int1, including 1450 bp upstream and 548 bp downstream from the Int1 open reading frame (www.candidagenome.org, Assembly 21), was integrated into the hisG locus of the int1-/- strain VBIDM2 (54) to produce the reconstituted strain KO509. Briefly, a copy of Int1 was generated by PCR, using primers 1 and 2 with BWP17 wt DNA as template and cloned into the SacI/MluI sites of pGEMHIS (51) to create pKO509. pKO509 was digested with SwaI and transformed into VBIDM2 to create the reconstituted strain KO509. PCR-mediated overlap extension mutagenesis (55) was used to produce copies of Int1 mutated at putative heparin binding domains. Briefly, primer pairs 1+3 and 2+4 (or 1+5 and 2+6) were used to produce two overlapping fragments of INT1 in which putative heparin binding domains were mutated (FKKRFFKL (SEQ ID NO: 2)→FKAAFFKL (SEQ ID NO: 53) or KQKKHQ (SEQ ID NO: 64)→KQAAHQ (SEQ ID NO: 63)), and a full length mutated sequence generated in a third per using primers 1+2 with the fragments as template. The mutated sequences were cloned into the SacI/MluI sites of pGEMHIS to create pKO503 and pKO507, respectively. A construct mutated at both sites (pKO508) was produced using primers 1+5 and 2+6 with plasmid DNA from pKO503 as template. Full length mutated products cloned into pGEMHIS and used to transform VBIDM2 as above, producing strains KO503, KO507, and KO508. The correct insertion and orientation of all constructs was confirmed by PCR. In growth curves performed with and without 100 units/ml heparin at 30° C. in RPMI-HEPES, there was no difference in the doubling times of the wild type, Δ int1 mutant, INT1 reintegrant, or the reintegrants containing alanine substitutions in Motif 1 (₈₀₄QKKHQIHK (SEQ ID NO: 1) to QAAHQIHK (SEQ ID NO: 52)), Motif 4 (₁₅₉₃FKKRFFKL (SEQ ID NO: 2) to FKAAFFKL (SEQ ID NO: 53)), or Motifs 1 and 4. Percent binding of each mutant to heparin in the ELISA assay was compared to the wild type according to the formula [Absorbance₅₉₅ mutant]/[Absorbance₅₉₅ wild type]×100.

The ELISA assay used a commercially available 96-well microtiter plate coated with polymerized allylamine (BD Biosciences) (27), in which wells were inoculated with 2.5

room temperature overnight, then washed with acetate buffer (100 mM NaCl, 50 mM NaOAc, 0.2% Tween, pH 7.2). Wells were washed three times with PBS, then inoculated with 1×10⁴ to 4×10⁴ *Candida albicans* yeast cells that were grown overnight at 30° C. in yeast peptone dextrose (YPD) broth, washed, and resuspended in RPMI-HEPES at 1×10⁷ organisms/ml. The plate was then incubated at 30° C. for one hour, then washed with PBS-Tween (PBST, PBS/0.05% Tween). A commercially available biotinylated anti-Candida antibody (Meridian Life Science, Memphis Tenn.) was diluted 1:1500 in FACS-TWEEN (0.3% BSA/0.05% Tween), added to the wells, incubated at 30° C. for one hour, then washed with PBST. Streptavidin-alkaline phosphatase (BioLegend, San Diego Calif.) was then added to the wells at a 1:10,000 dilution in FACS-TWEEN and incubated at 30° C. for 30 min, then washed with PBST and AKP buffer (100 mM Tris, 100 mM NaCl, 50 mM MgCl₂·6H₂O). BluePhos[®] microwell substrate (Kirkegaard and Perry Laboratories, Gaithersburg Md.) was added to the wells and allowed to react for 45 min at room temperature. The color change was read spectrophotometrically at absorbance of 595 nm.

FIGS. 4A-D show that linear heparin binding motifs in *C. albicans* Int1 mediate binding to heparin.

FIG. 4A illustrates the five putative heparin binding sites in Int1. Mutations were made by substituting alanine residues in Motif 1: ₈₀₄QKKHQIHK (SEQ ID NO: 1) to QAAHQIHK (SEQ ID NO: 52) (Table 1, Motif 1) and in Motif 4: ₁₅₉₃FKKRFFKL (SEQ ID NO: 2) to FKAAFFKL (SEQ ID NO: 53) (Table 1, Motif 4). A third reintegrant had alanine substitutions at both sites.

FIG. 4B shows molecular modeling of Motif 4, which predicts that the three lysine residues and the single arginine residue (arrows) might facilitate binding to a strong anion like heparin.

FIG. 4C shows heparin binding of *C. albicans* WT (normalized to 100%), Δ int1 double disruptant (DD), and four single-copy reintegrants: (1) Reint contains one wild type copy of Int1; (2) Motif 1 mutant contains alanine substitutions for lysine residues at positions 805/806 (₈₀₄QAAHQIHK (SEQ ID NO: 52)); (3) Motif 4 mutant contains alanine

substitutions for lysine 1595 and arginine 1596 (1593FKAAFFKL (SEQ ID NO: 53)), and (4) Motif 1&4 mutant contains alanine substitutions at both sites. Binding to heparin was highest for the wild-type *C. albicans* strain that expressed both copies of the INT1 gene. Disruption of both copies of INT1 (DD) reduced the binding of heparin by 40% ($p=0.031$) despite removing just one of the 34 *C. albicans* genes encoding putative linear HBM. Reintegration of one copy of the wild type INT1 gene (Reint) partially restored the ability of *C. albicans* to bind to heparin. Differences in heparin binding among the WT, Reint and DD strains were statistically significant ($p=0.031$ in all cases). Among the three isogenic strains with alanine mutations in putative heparin binding sites, mutation of Motif 1 was associated with the largest reduction in heparin binding, compared to the wild type strain ($p=0.031$), to the Motif 4 mutant ($p=0.036$), and to the Motif 1&4 mutant ($p=0.031$). Results with the Motif 1 mutant were not significantly different from the percent binding observed with the double disruptant ($p=0.063$) or the reintegrand ($p=0.115$), nevertheless, this trend of reduced binding suggested that lysine residues in Motif 1 might be likely mediators of heparin binding. Results are presented as mean \pm SD for $n=5$ experiments performed in quadruplicate, * $p=0.031$, ** $p=0.036$.

Because Motif 1 in Int1 appeared to mediate a considerable proportion of the binding of *C. albicans* to heparin, a peptide encompassing this motif HKQEKQKKHQIHKV (SEQ ID NO: 4) was used to immunize rabbits with a commercial protocol (Pacific Immunology). Affinity-purified immune IgG and IgG from pre-immune rabbit serum (both from Pacific Immunology) were compared for the ability to inhibit binding of *C. albicans* to heparin (FIG. 5). Antibody and heparin inhibition studies were performed as follows. Antibody was produced and tested by Pacific Immunology, Ramona, Calif. (www.pacificimmunology.com; NIH Animal Welfare Assurance Number A41820-01; USDA License 93-R-283). A peptide corresponding to amino acids 799-812 of Int1 (HKQEKQKKHQIHKV (SEQ ID NO: 4)), encompassing Motif 1, was conjugated to KLH via an N-terminal cysteine residue and used to immunize two NZW rabbits. The animals were immunized once with conjugated peptide in a proprietary formulation of Freund's complete adjuvant and boosted 3 times with conjugated peptide in Freund's incomplete adjuvant. The same peptide was conjugated to CNBr-Sepharose and used for affinity purification of epitope-specific IgG. The final serum titer for both animals was $>1:100,000$ by ELISA. Both pre-immune rabbit serum and serum from bleed 3 were chromatographed on a 2 ml Protein A column (Thermo Scientific) to yield pre- and post-immune IgG and brought to a concentration of 1.6 mg/ml. 100 μ l poly-D-lysine (MP Biomedicals, Solon Ohio) diluted as 1 mg in 10 ml distilled water was added to each well of black 96-well plates (Costar) and incubated at room temperature for 60 min. After three washes with PBS, 100 μ l pharmacologic heparin (2000 units/ml in RPMI-HEPES) was added to each well and incubated overnight at room temperature in the dark. In the morning, the plate was washed three times with PBS, then a 1:1000 dilution of pre- or post-immune IgG in PBS was added to each well and incubated for 30 min at room temperature. After overnight growth to mid-exponential phase in CSM, *C. albicans* wild type and double disruptant strains were labeled with CFSE as above. After labeling, samples of each strain were removed to determine the intensity of labeling by flow cytometry, as described above. CFSE-labeled *C. albicans* were diluted in RPMI-HEPES to yield 1×10^6 cells per 100 μ l and added to each well for 60 min at 30° C. The plate was washed three times with PBS; fluorescence in each

well was measured with a Beckman DTS 880. Experiments were performed in triplicate. For heparin inhibition studies, wild type *C. albicans* were grown as above, labeled with CFSE, and washed to remove excess CFSE. Organisms were pelleted, the supernatant was removed, and the pellets were suspended in 2 ml pharmacologic heparin (20,000 units/ml) diluted with RPMI-HEPES to a concentration of 250 units/ml (1.3 mg/ml). CFSE-labeled *C. albicans* were incubated with heparin for 60 min at 37° C. with shaking at 400 RPM, then added to the wells of a black plate, incubated for 60 min at RT, and washed three times with PBS; fluorescence in each well was measured with a Beckman DTS 880. The assay was performed in a 96-well black plate with CFSE-labeled *C. albicans*. Wells were coated with poly-D-lysine, 100 μ g/100 μ l PBS for 60 min at room temperature, then 200 units heparin in 50 μ l RPMI-HEPES were added to each well. The plate was incubated overnight at room temperature. The following afternoon, pre-immune IgG and post-immune IgG were diluted in PBS to a final concentration of 13.2 μ g/ml. 50 μ l of pre- or post-immune IgG was deposited in the appropriate well for 30 min at room temperature, then 1×10^6 CFSE-labeled *C. albicans* in 50 μ l RPMI-HEPES were added for two hours at room temperature. After this incubation, the wells were washed three times with PBS, and fluorescence was measured on the multimode detector as previously described.

Polyclonal IgG to the HKQEKQKKHQIHKV (SEQ ID NO: 4) motif in domain 3 of Int1 blocked 19% of binding of wild type *C. albicans* (WT) to heparin. FIG. 5 shows heparin binding of wild type *C. albicans* preincubated with pre-immune IgG (left closed circles) and wild type *C. albicans* pre-incubated with post-immune IgG raised to the polypeptide HKQEKQKKHQIHKV (SEQ ID NO: 4) (right closed circles), where data represent mean \pm SD of $n=15$ experiments performed in duplicate, * $p=0.023$, by paired statistics. These results confirmed the importance of Motif 1, as a mediator of heparin binding in *C. albicans*.

In other systems, heparin has been reported to cleave surface proteins (56), change protein conformation (57), and bind trypsin-sensitive lysine and arginine residues in histone H1 (58). In order to understand whether incubation with heparin changed the conformation of *C. albicans* surface proteins, 1×10^7 *C. albicans* were incubated for one hour at 3° C. on a rotator with pharmacologic heparin (20,000 units/ml) diluted in RPMI-HEPES to a working concentration of 250 units/ml. *C. albicans* 1×10^7 *C. albicans* in an equal volume of RPMI-HEPES served as control. Supernatants were removed. 300 μ l of Hep+ and Hep- supernatants were incubated with 100 μ l avidin agarose beads (Thermo Scientific) for 60 min at room temperature on a rotator in the presence of 50 units heparin (Hep+); an equal volume of RPMI-HEPES was substituted for heparin in the Hep- supernatants. Beads were pelleted, and 100 μ l beads were incubated with 100 μ l 3.0 M NaCl for 30 mins at room temperature on a rotator. Beads were pelleted and the supernatants were withdrawn; 100 μ l TCA was added to 100 μ l Hep+ or 100 μ l Hep- supernatants and incubated on ice overnight at 4° C. The TCA precipitates were stored at -80° C. until analysis by mass spectroscopy.

Six biological replicates of TCA precipitated protein pellets from equal cellular equivalence of heparin-treated (Hep+) and untreated (Hep-) conditioned medium from cultures of *C. albicans* were each solubilized in 50 μ l Laemmli gel buffer. To remove any residual TCA and to further concentrate the samples, each was subjected to buffer exchange and concentration using an Amicon ultra 3 kDa microfuge filtration cartridge at 14,000 \times g for 15 min with five subse-

quent additions of 50 μ L of 1 \times Laemmli gel buffer between spins. The resulting retained proteins (6 Hep+ and 6 Hep-) were subsequently prepared for SDS-PAGE by combining into two pools of three samples for the Hep+ and Hep- conditioned medium. The replicate sample pools were loaded onto two 4-12% mini gels and separated using the MOPS buffer system followed by silver staining to visualize the proteins using the Sigma Proteosilver system. The proteins were prepared for identification and quantitation by mass spectrometry by gridding the gel lanes from the replicates of Hep+ and Hep- samples into 11 equal regions followed by in gel trypsin digestion and extraction of peptides as described (59)

The recovered peptides from the gridded gel sections were analyzed by liquid chromatography coupled nano-electrospray mass spectrometry (nLC-MS/MS) on a TripleTOF 5600 mass spectrometer (AB Sciex, Toronto Canada) attached to an Eksigent (Dublin Calif.) nanoLC ultra nanoflow system. Recovered peptides from each fraction were loaded on to IntegraFrit Trap Column (outer diameter of 360 μ m, inner diameter of 100, and 25 μ m packed bed) from New Objective, Inc. (Woburn Mass.) at 2 μ L/min in FA/H₂O 0.4/99.2 (v/v) for 10 min to desalt and concentrate the samples. For the chromatographic separation of peptides, the trap-column was switched to align with the analytical column, Acclaim Pep-Map100 (inner diameter of 75 μ m, length of 15 cm, C18 particle sizes of 3 μ m and pore sizes of 100 Å) from Dionex-Thermo Fisher Scientific (Sunnyvale Calif.). The peptides were eluted using a linear gradient from 95% phase A (FA/H₂O 0.4/99.6, v/v) to 40% phase B (FA/ACN 0.4/99.6, v/v) from 5 min to 22.5 min (2% ACN/min) at a flow of 300 mL/min. As the peptides eluted from the column they were sprayed into the mass spectrometer using NANOSpray® III Source (AB Sciex, Toronto Canada). Ion source gas 1 (GS1), ion source gas 2 (GS2) and curtain gas (CUR) were respectively kept at 7, 0 and 25 vendor specified arbitrary units. Interface heater temperature and ion spray voltage was kept at 150 C, and at 2.3 kV respectively. Mass spectrometer method was operated in positive ion mode set to go through 25 minutes, where each cycle performing one TOF-MS scan type (0.25 sec accumulation time, in a 400 to 1600 m/z window) followed by twenty information dependent acquisition (IDA)-mode MS/MS-scans on the most intense candidate ions selected from initially performed TOF-MS scan during each cycle having a minimum 150 counts. Each product ion scan was operated under vendor specified high-sensitivity mode with an accumulation time of 0.05 sec and a mass tolerance of 50 mDa. Former MS/MS-analyzed candidate ions were excluded for 10 sec after its first occurrence, and data were recorded using Analyst®-TF (1.5.1) software.

NanoLC-MS/MS data collected from the Hep+ and Hep- samples were converted to Mascot generic files and searched against the SwissProt fungal database on an in house server running Mascot version 2.2.07 (Matrix Science, Ltd). Specific search parameters included up to two missed tryptic cleavages, carbamidomethylation of Cys, oxidation of Met, peptide and fragmentation mass tolerance of 0.1 Da. Only proteins with a minimum of two peptides with Mascot peptide score indicating a peptide identity and a false discovery rate (FDR) against and inverse database at less than 1% were reported. Semi-quantitative measurements between the Hep+ and Hep- proteins were generated using a minimum of two tryptic peptides from each protein as surrogates for the amount of proteins from the two groups. This was accomplished by capturing extracted ion profiles for each peptide and then comparing the mono-isotopic peak intensity at the apex of the signal for the M+2H or M+3H signal for each

peptide. An average ratio of the maximal peak intensity between two independent peptides for each protein presented was used to determine semi-quantitative ratios of proteins between the Hep+ and Hep- samples.

To determine the levels of these proteins in control versus heparin-treated organisms, the relative level of several tryptic peptides were used as surrogates for the protein levels.

Compared to untreated organisms, heparin treatment led to a seven-fold increase in intensity of IDVVDQAK (SEQ ID NO: 65) from Eno1 (FIG. 6A), a ten-fold increase in intensity of SLLDAAVK (SEQ ID NO: 66) from Pgk1 (FIG. 6B), and a five-fold increase in VPTTDSVVDLTVR (SEQ ID NO: 67) from Tdh3 (FIG. 6C) in supernatants.

Other peptides from Eno1, Pgk1, and Tdh3 also exhibited increased intensities, ranging from 10-16 fold the intensity of the corresponding peptides from untreated organisms (data not shown), further validating significant increases in the protein levels in the supernatants for heparin-treated cultures. Of the 12 proteins whose peptides were found in highest concentration in the supernatant (Table 4), all but one are known to be localized to the cell wall; cellular localization of Eft1 is not known. Eight of the twelve, including Eno1, Pgk1, Tdh3 and Ssa1/2, themselves contain putative heparin binding motifs (Table 1) and are considered critical antigens for innate and adaptive immune responses against *C. albicans* (31-36).

TABLE 4

ID	Prot Score	Prot Mass	Peptide Matches**
Eno1* [#]	2877	47202	213
Ssa2* [#]	1938	70199	73
Pgk1* [#]	1826	45266	131
Tdh3* [#]	1751	35508	192
Hsp90 [#]	1666	80773	65
Ssb1* [#]	1608	66562	82
Met6 [#]	1566	85763	70
Ssa1* [#]	1455	70452	61
Adh1 [#]	1218	37255	67
Eft1	1187	50426	99
Ino1* [#]	1006	57857	39
Eft2* [#]	933	93865	25

*proteins with putative linear heparin binding motifs;

**total peptides identified by MS/MS including duplicates; and

[#]cell wall proteins by Gene Ontology.

In order to evaluate the consequences of removal of Ssa1 and Ssa2 (Table 4), both targets of histatin, a histatin killing assay was performed as follows. *C. albicans* strains were grown to late log phase overnight in YPD. A single colony was suspended in 1 ml YPD, diluted 1:500 into 10 ml YPD, and incubated overnight at 30° C. and 225 rpm to OD₆₀₀ 1.0. Yeast cells were washed twice in PBS and 2 \times 10⁴ cells suspended in 250 μ L RPMI+10 mM HEPES, pH 7.0, with or without 500 units/ml heparin (Sigma). The cells were incubated at 37° C. for one hour with shaking, washed twice with 10 mM phosphate buffer, pH 7.4, and suspended in 20 μ L phosphate buffer. Histatin 5 (final concentration 15 μ M; Peptides International) or 10 mM phosphate buffer was added to the preincubated cells (total volume 40 μ L) and incubated further at 37° C. for 90 minutes with shaking. YPD (360 μ L) was added to each tube, a 40 μ L aliquot spread onto YPD plates, and colonies counted after two days. The effect of heparin treatment was determined by the formula % change = [(cfu+heparin)-(cfu-heparin)]/(cfu+heparin).

Heparin binding to trypsin-sensitive lysine and arginine residues in histone H1 unfolds chromatin and increases its accessibility (58). To test whether heparin binding could influence gene expression in *C. albicans*, qRT-PCR was per-

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formed on a selective set of 13 genes involved in adhesion, cell-cell interaction, and biofilm formation after incubating *C. albicans* with and without 100 units/ml heparin at 37° C. for 75 minutes. Gene expression studies were performed as follows. Overnight cultures (3 ml YPD at 30° C., 225 rpm) were diluted in 25 ml YPD to a nominal concentration of 8×10^5 /ml and grown to OD₆₀₀ approximately 1.0 (30° C., 225 rpm), collected by centrifugation, and washed in PBS. Cells (5×10^7) were suspended in 5 ml RPMI1640 with 25 mM MOPS pH7.4, with or without 100 units/ml Sigma heparin in 50 ml polypropylene tubes, and incubated at 37° C. with shaking for 75 minutes. One ml aliquots (approximately 10^7 cells) were collected by centrifugation, washed once with room temperature PBS, and frozen at -80°. Pellets were thawed on ice and suspended in 1 ml Tri-Reagent (MRC Research) in a 2 ml screw capped tube containing about 0.2 g acid washed glass beads (Sigma) and vortexed three times for 1 minute with a 1 minute rest on ice between each vortex. The lysates were rested for five min at room temperature, centrifuged for five min at 12,000 rpm, and RNA isolated from the supernatant using the DirectZol kit (Zymo), including DNase digestion, per manufacturer's instructions. cDNA was produced from equivalent quantities of RNA (between 300 and 900 ng) for each treatment using the Maxima Reverse Transcriptase Kit for qRT-PCR (Fermentas). Two µl of 1:5 dilution of cDNA was used in each qPCR reaction with 500 nM of each primer and Fast SYBR Green Master Mix (Life Technologies) on a 7500 FAST Instrument (Applied Biosystems) according to manufacturer's instructions. Relative expression was determined using the $\Delta\Delta C_t$ method (60) with 18S RNA as the reference. Primers are shown in Table 5.

TABLE 5

Primer	Sequence	SEQ ID NO:
18S F	TCTTGTGAACTCCGTCGTG	68
18S R	AGGGACGTAATCAACGCAAG	69
AHP1 F	TGTGCTGGTGCTTTTACC	70
AHP1 R	TTAGCCCAAGCTGCCATTAC	71
ALS1 F	TCATTTGCCACCACTACCAC	72
ALS1 R	TGGCATAGGATTGTGACCAG	73
ALS3b F	GCTGGTGGTTATTGGCAACGTGC	74
ALS3b R	TGGTAAGGTGGTCACGGCGG	75
CDC10 F	AGATCAAGGGCAAACCTCAC	76
CDC10 R	ATAGGAGCATTTGGCACACC	77
EAP1 F	TACCCAGGCCAATACAAAGG	78
EAP1 R	TAATGGGCTTGACCTTGGAG	79
ECE1 F	CTAATGCCCTCGTCAGATTG	80
ECE1 R	AACATCTGGAACGCCATCTC	81
ENO1 F	CCATTGACAAAGCCGGTTAC	82
ENO1 R	TTAGATGGGTGGGATTCTGG	83
HGC1 F	AGGTCGCAAGCAACAACAAC	84
HGC1 R	AAGAAACAGCACGAGAACCCAG	85
HWP1b F	TCCTGCCACTGAACCTTCCCCAG	86

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TABLE 5 -continued

Primer	Sequence	SEQ ID NO:
HWP1b R	CCACTTGAGCCAGCTGGAGCG	87
HWP2 F	CCACCAAACCAAGTGCTAC	88
HWP2 R	AACTCCAGATGATCCCGAAG	89
INT1 F	TGTGCCCCACTGAAGTCAAAG	90
INT1 R	GCTTTACCGGTGATTGGTC	91
RBT1 F	CACCTCATGCTCCAACAATG	92
RBT1 R	GATGATTCTGGGGCTGATTC	93
RBT5 F	TGCTGAAAGTTCTGCACCAG	94
RBT5 R	GCTTCAACGGAACAGAAGC	95

FIG. 7A shows CFU of *C. albicans* WT and DD after 75-minute incubation without (left bar of each set) or with (right bar of each set) 500 units/ml heparin followed by histatin 5 (15 µM), performed in duplicate. Confirming proteomic results showing removal of Ssa1/2 (Table 3), incubation of wild type *C. albicans* with 500 units/ml heparin led to a 25% decrease in histatin-mediated killing (FIG. 7A); in contrast, heparin treatment of the INT1 double disruptant did not impair killing (1.5% decrease).

FIG. 7B shows relative mRNA expression of thirteen genes measured by qRT-PCR after incubation of *C. albicans* with heparin, with the level of expression compared to *C. albicans* without heparin (=1). Results showed a 2.5 to 3-fold increase in mRNA for HWP1 and HWP2.

In order to understand whether heparin binding motifs 1 and 4 influenced *C. albicans* pathogenesis in vivo, we employed a rat model of biofilm formation in central venous catheters inserted into the jugular vein, as previously described (61). After insertion into the jugular vein of an anesthetized female Sprague-Dawley rat, a silastic catheter was heparinized with 100 units heparin/ml and remained in place for 24 hours. At the 24 hour timepoint, 500 µl of blood was withdrawn and cultures to insure sterility, and then 1×10^6 CFU of the desired *C. albicans* strain was instilled into the catheter and allowed to dwell for 6 hours. The animal was then sacrificed, the catheter was removed aseptically and processed for scanning electron microscopy. Biofilm formation on the intra-luminal surface of the catheter was assessed by scanning electron microscopy (SEM) at 100× and 2000×.

FIGS. 8A-F demonstrated that heparin binding motifs contribute to biofilm formation in vivo.

The INT1 wild type strain (WT) produced a profuse biofilm with intertwined hyphae and visible exopolysaccharide matrix (FIG. 8A). Biofilm formation by the Int1 double disruptant was much reduced on SEM (FIG. 8B), as expected. Reintegration of one wild type copy of INT1 restored a profuse biofilm (FIG. 8C). However, alanine substitution of lysines_{805/806} in Motif 1 greatly impaired biofilm formation (FIG. 8D). Although alanine substitution of lysine₁₅₉₅ and arginine₁₅₉₆ in Motif 4 did not reduce biofilm formation (FIG. 8E), the Motif 1&4 mutant again produced sparse biofilm (FIG. 8F). These results showed that lysine residues 805/806 in Motif 1 were critical for biofilm formation in vivo.

Inhibition of biofilm formation by the antibody raised against the peptide sequence HKQEKQKKHQIHKV (SEQ ID NO:4) was also tested in the rat central venous catheter model. A 1:10 dilution of affinity-purified IgG against the peptide HKQEKQKKHQIHKV (SEQ ID NO: 4) was incu-

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bated with wild type *C. albicans* at 30° C. for one hour. A 1:10 dilution of pre-immune IgG was used as a control. The strains were then instilled into separate jugular venous catheters in individual rats. After 6 hours, catheters were removed and aseptically processed for scanning electron microscopy (100× and 2000×) as described above.

Central venous catheters from animals that received *C. albicans* pre-incubated with IgG against HKQEKQKKHQIHKV (SEQ ID NO:4) exhibited substantially reduced biofilm formation. FIG. 9 (left panel) shows intraluminal biofilm (100× and 2000×) from *C. albicans* incubated with pre-immune IgG; there is no diminution in biofilm, hyphae, or production of exopolysaccharide matrix. FIG. 9 (right panel) shows intraluminal biofilm (100× and 2000×) from *C. albicans* incubated with post-immune IgG recognizing the sequence HKQEKQKKHQIHKV (SEQ ID NO: 4); there is a marked diminution in biofilm with sparse hyphae and no matrix.

In vitro study results showed that *C. albicans* binds heparin through HBM in Int1 (FIG. 4C). The specificity of this interaction was confirmed by inhibition with heparin (data not shown) and with antibodies directed against a peptide that encompasses Motif 1 in Int1 (FIG. 5). Binding of heparin results in several consequences that could potentially impact virulence in vivo: removal of *Candida* surface proteins that serve as targets for innate (FIG. 7A) and adaptive (FIG. 6A-C) immune defenses and modulation of gene expression (FIG. 7B).

The in vivo studies of biofilm formation in heparinized central venous catheters in rats showed an obvious reduction in biofilm formation after mutation of lysine residues 805/806 in Motif 1 (FIGS. 8D, 8F). In addition, a rabbit IgG antibody directed against a peptide encompassing Motif 1 dramatically inhibited biofilm formation in the rat central venous catheter model as well (FIG. 9, right panel). These results not only demonstrate the central role of lysine residues in Motif 1 but also have important clinical implications because of the use of heparin in central venous catheters, in which setting *Candida* spp. are the fourth most common cause of infections (23, 24).

Putative linear HBM are also present in *Staphylococcus epidermidis* and *Staphylococcus aureus* (Long and Hostetter unpublished data), two organisms that are even more common causes of catheter-associated infection (43), as shown in Tables I-IV. For example, putative linear HBM were identified

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in the following cell wall or putative cell wall proteins, where the motif and the beginning position of the motif is indicated in parenthesis, from methicillin resistant *Staphylococcus aureus*, strain 252 (MRSA252): sasC (LKKNKY; 4; and IRKYKV; 11), isdB (YKKAKT; 289), sasF (SRRNKL; 618), glcB (IRKFKL; 415), sasA (MHHTHS; 1263), SAR0879 (LKKIKG; 573), SAR0986 (YRHLKP; 754), SAR1559 (IRKAHQ; 206), SAR2393 (PKRKVVKI; 149), and sasG (VRKARS; 140); from methicillin sensitive *Staphylococcus aureus*, strain 476 (MSSA476): SAS2383 (VRKARS; 140; and VKKSKI; 1319), SAS1682 (LKKNKY; 4; and IRKYKV; 11), SAS1063 (YKKAKT; 282), SAS1657 (YHKAKT; 484), SAS2532 (SRRNKL; 626), SAS2540 (MHHTHS; 2187), SAS0082 (LKKIKG; 573), SAS1011 (IRKAHQ; 206), SAS2035 (PKRKVVKI; 149), and SAS2424 (IRKFKL; 415); and from *Staphylococcus epidermidis*, strain RP62A: SERP1316 (FRKQKF; 4; VHRLKV; 352; and IHKIKP; 3234), SERP0660 (LKKWKV; 4; and IRRAHQ; 212), SERP0719 (TRKNHY; 13), SERP1482 (VKRFKN; 1730), SERP1654 (MKKSKV; 1), SERP2264 (MKRIKT; 393), SERP0207 (NRKNKN; 887), and SERP1691 (PKKIKN; 72).

In one embodiment, an antibody is generated against a linear heparin binding motif which is conserved among MSSA, MRSA, and *S. epidermidis*. In one embodiment, the conserved heparin binding motif is selected from the group consisting of LKKNKY, LKKNKY, LKKWKV, LKKIKG, LKKIKG, VRKARS, VRKARS, YKKAKT, YKKAKT, YHKAKT, PKRKVVKI, PKRKVVKI, IRKAHQ, IRKAHQ, and IRRAHQ.

Putative linear HBM were identified in the following cell wall or putative cell wall proteins from various yeast species, as shown in Table V where a check mark indicates that the motif is identical to the motif found in *C. albicans*, including *C. dubliniensis* (Int1 (YKKRFFKL), Eno1 (AKKGKF), and Tdh3 (GHKIKV) proteins), *C. parapsilosis* (Tdh3 protein (GHKIKV)), *C. tropicalis* (Int1 (FKRRFFKL), and Tdh3 (GHKIKV) proteins), *C. glabrata* (Int1 (FKRRFFTL) protein), *Lodderomyces elongisporus* (Int1 (FKKFIFKL) and Tdh3 (GHKIKV) proteins), and *A. nidulans* (Int1 (FKRRFFKL) protein). In embodiments, an antibody directed to a region of these proteins containing the putative HBM or the region of these proteins containing the putative HBM may be used in the described methods.

TABLE I

Methicillin resistant <i>Staphylococcus aureus</i> , strain 252 MRSA252									
GI	Accession	Name	Description	"Cell wall" in Description	"Cell wall" in GO annotation	Have signaling peptide	Motif in Se- quence	Motif	Start Loca- tion
GI:49484003	sasC		putative surface anchored protein			Yes	Yes	LPNTG	2153
GI:49483291	isdB		iron-regulated heme-iron binding protein			Yes	Yes	LPQTG	616
GI:49484843	sasF		putative surface anchored protein			Yes	Yes	LPKAG	588
GI:49484739	glcB		PTS system, glucose-specific IIA _{BC} component				Yes	LPAAG	22
GI:49484850	sasA		putative serine rich repeat containing protein				Yes	LPDTG	1308
GI:49482351	SAR0879		putative myosin-crossreactive antigen				Yes	LPKAG	57
GI:49482500	SAR0986		putative nitric oxide reductase				Yes	LPSAG	230

TABLE I -continued

Methicillin resistant <i>Staphylococcus aureus</i> , strain 252 MRSA252						
GI:49483239	SAR1559	putative cobalt transport protein			Yes	LPITG 258
GI:49484356	SAR2393 sasG	hypothetical protein virulence associated cell wall protein			Yes	LPTAG 177
GI Accession	Ortholog in MSSA476	ortholog has motif?	Ortholog in RP62A	RP62A orthology has motif:	Have heparin binding motif?	#of heparin binding motifs
GI:49484003	SAS1682	Yes	SERP1316	Yes	Yes	2
GI:49483291	SAS1063	Yes			Yes	1
GI:49484843	SAS2532	Yes	SERP2264	Yes	Yes	1
GI:49484739	SAS2424	Yes			Yes	1
GI:49484850	SAS2540	Yes	GI:57865710	No	Yes	1
GI:49482351	SAS0082	Yes	GI:57866574	No	Yes	1
GI:49482500					Yes	1
GI:49483239	SAS1011	Yes	SERP0660	Yes	Yes	1
GI:49484356	SAS2035	Yes	SERP1739	Yes	Yes	1
					Yes	1
GI Accession	Motif 1 type	Motif 1 seq	Motif after	Motif 2 type	Motif 2 seq	Motif lifter
GI:49484003	Cardin	LKKNKY	3	Cardin	IRKYKV	10
GI:49483291	Cardin	YKKAKT	288			
GI:49484843	Cardin	SRRNKL	617			
GI:49484739	Cardin	IRKFKL	414			
GI:49484850	Cardin	MHHTHS	1262			
GI:49482351	Cardin	LKKIKG	572			
GI:49482500	Cardin	YRHLKP	753			
GI:49483239	Cardin	IRKAHQ	205			
GI:49484356	Wentraub	PKRKVVKI	148			
	Cardin	VRKARS	139	found by MKH		

TABLE II

Methicillin sensitive <i>Staphylococcus aureus</i> , strain 476 MSSA476						
ID MSSA476	refSeq	GI Accession	Name	Description	"Cell wall" in Description	"Cell wall" in GO annotation
SAS2383	YP_044496.1	GI:49487275	SAS2383	putative cell wall-anchored protein	Yes	Yes
SAS1682	YP_043802.1	GI:49486581	SAS1682	putative surface anchored protein		
SAS1063	YP_043187.1	GI:49485966	SAS1063	iron-regulated heme-iron binding protein		Yes
SAS1657	YP_043776.1	GI:49486555	SAS1657	haptoglobin-binding surface anchored protein		Yes
SAS2532	YP_044646.1	GI:49487425	SAS2532	putative surface anchored protein		
SAS2540	YP_044654.1	GI:49487433	SAS2540	putative cell wall-anchored protein	Yes	Yes
SAS0082	YP_042206.1	GI:49484985	SAS0082	putative myosin-crossreactive antigen		
SAS1011	YP_043135.1	GI:49485914	SAS1011	putative cobalt transport protein		
SAS2035	YP_044146.1	GI:49486925	SAS2035	hypothetical protein		
SAS2424	YP_044538.1	GI:49487317	SAS2424	PTS system, glucose-specific IIABC component		

TABLE II -continued

Methicillin sensitive <i>Staphylococcus aureus</i> , strain 476 MSSA476									
ID MSSA476	Have signaling peptide	Motif in Sequence	Motif	Start Loca- tion	Ortholog in MSSA476	MRSA252 ortholog has motif?	Ortholog in RP62A	RP62A orthology has motif?	Have heparin binding motif?
SAS2383	Yes	Yes	LPKTG	1338	sasG		aap	Yes	Yes
SAS1682	Yes	Yes	LPNTG	2150	sasC	Yes	SERP1316	Yes	Yes
SAS1063	Yes	Yes	LPQTG	609	isdB	Yes			Yes
SAS1657	Yes	Yes	LPKTG	860					Yes
SAS2532	Yes	Yes	LPKAG	596	sasF	Yes	SERP2264	Yes	Yes
SAS2540		Yes	LPDTG	2232	sasA	Yes	GI:57865710	No	Yes
SAS0082		Yes	LPKAG	57	SAR0879	Yes	GI:57866574	No	Yes
SAS1011		Yes	LPITG	258	SAR1559	Yes	SERP0660	Yes	Yes
SAS2035		Yes	LPTAG	177	SAR2393	Yes	SERP1739	Yes	Yes
SAS2424		Yes	LPAAG	22	glcB	Yes			Yes

ID MSSA476	# of heparin binding motifs	Motif 1 type	Motif 1 seq	Motif after	Motif 2 type	Motif 2 seq	Motif after
SAS2383	2	Cardin	VRKARS	139	Cardin	VKSKI	1318
SAS1682	2	Cardin	LKKNKY	3	Cardin	IRKYKV	10
SAS1063	1	Cardin	YKKAKT	281			
SAS1657	1	Cardin	YHKAKT	483			
SAS2532	1	Cardin	SRRNKL	625			
SAS2540	1	Cardin	MHHTHS	2186			
SAS0082	1	Cardin	LKKIKG	572			
SAS1011	1	Cardin	IRKAHQ	205			
SAS2035	1	Wentraub	PKRKVVKI	148			
SAS2424	1	Cardin	IRKFKL	414			

TABLE III

Staphylococcus epidermidis, strain RP62A											
ID	RP62A	refSeq	GI	Accession	Name	Description		"Cell wall" in Description	"Cell wall" in GO annotation		
SERP1316	YP_188888.1	GI:57867198	SERP1316	cell wall surface anchor family protein				Yes			
SERP0660	YP_188245.1	GI:57866567	SERP0660	cobalt transport family protein							
SERP0719	YP_188302.1	GI:57866639	SERP0719	cell wall surface anchor family protein				Yes			
SERP1482	YP_189048.1	GI:57867352	SERP1482	cell wall surface anchor family protein				Yes			
SERP1654	YP_189219.1	GI:57867536	SERP1654	cell wall surface anchor family protein				Yes			
SERP2264	YP_189815.1	GI:57865679	SERP2264	cell wall surface anchor family protein				Yes			
SERP0207	YP_187803.1	GI:57866125	SERP0207	sdrG protein							
SERP1691	YP_189256.1	GI:57867615	SERP1691	cell division protein, FtsW/RodA/SpoVE family							
Strain RP62A (continuation)											
ID	RP62A	Have sig- naling peptide	Motif in Se- quence	Motif	Start Loca- tion	Ortholog in MSSA476	MSSA476 ortholog has motif?	Ortholog in MRSA252	MRSA252 orthology has motif?	Have heparin binding motif?	# of heparin binding motifs
SERP1316	Yes	Yes	LPEAG	3654	SAS1682	Yes	sasC	Yes	Yes	Yes	3
SERP0660		Yes	LPITG	264	SAS1011	Yes	SAR1559	Yes	Yes	Yes	2

TABLE III -continued

<i>Staphylococcus epidermidis</i> , strain RP62A									
SERP0719	Yes	Yes	LPETG	787				Yes	1
SERP1482	Yes	Yes	LPDTG	1937				Yes	1
SERP1654	Yes	Yes	LPETG	165				Yes	1
SERP2264	Yes	Yes	LPATG	639	SAS2532	Yes	sasF	Yes	1
SERP0207		Yes	LPDTG	852	SAS0521	Yes	bbp	Yes	1
SERP1691		Yes	LPITG	352	SAS1988	Yes	SAR2371	Yes	1

Strain RP62A (continuation)									
ID RP62A	Motif 1 type	Motif 1 seq	Motif after	Motif 2 type	Motif 2 seq	Motif after	Motif 3 type	Motif 3 seq	Motif after
SERP1316	Cardin	FRKQKF	3	Cardin	VHRLKV	351	Cardin	IHKIKP	3233
SERP0660	Cardin	LKKWKV	3	Cardin	IRRAHQ	211			
SERP0719	Cardin	TRKNHY	12						
SERP1482	Cardin	VKRFGN	1729						
SERP1654	Cardin	MKSKV	0						
SERP2264	Cardin	MKRIKT	392						
SERP0207	Cardin	NRKNKN	886						
SERP1691	Cardin	PKKIKN	71						

TABLE IV

Linear Heparin Binding Motifs Conserved Among 2 or 3 <i>Staphylococcus</i> species									
MRSA	sasC	LKKNKY	sar0879	LKKIKG	sasG	VRKARS	isdB	YKKAQT	sar2892
MSSA	sasC	LKKNNY	sas0082	LKKIKG	sas2383	VRKARS	sas1063	YKKAQT	sas1657
St. epi	serp0660	LKKWKV						YHKAQT	sas2035

MRSA	PKRKVVKI	sari 559	IRKAHQ
MSSA	PKRKVVKI	sas1011	IRKAHQ
St. epi		serp0660	IRKAHQ

TABLE V

Heparin Binding Motifs in various yeast species								
HEPARIN-BINDING MOTIF								
<i>C. albicans</i> XXXXXXXX	<i>C. albicans</i>	<i>C. dubliniensis</i>	<i>C. parapsilosis</i>	<i>C. tropicalis</i>	<i>C. glabrata</i>	<i>Lodderomyces elongisporus</i>	<i>A. nidulans</i>	
Motif	Protein							
FKKRFFKL	Int1	✓	YKKRFFKL	no	FKRRFFKL	FKKRFFTL	FKKFIFKL	✓
LRLRLT	Ssa2/Ssb1	✓					no	
AKKGKF	Eno1	✓	✓	no	no		no	
GHIKIV	Tdh3	✓	✓	✓	✓		✓	

✓ identical

The interaction of *C. albicans* with heparin or heparin-like compounds may play a key role in clinical settings where heparin is used (e.g. central venous catheters, dialysis catheters) or possibly in tissues where heparan sulfates are preferentially expressed.

Because antibodies raised against a linear heparin binding motif inhibited binding of *C. albicans* to heparin, the heparin binding peptide HKQEKQKKHQIHKV (SEQ ID NO: 4) could be used as an immunizing antigen in immunocompetent patients to elicit antibodies that protect against heparin binding.

For example, a peptide from streptococcal M protein has been used to make a vaccine for rheumatic fever (62). Alternatively, antibodies raised against this peptide in immunized humans can be used for passive immunization. This technology is used with commercially available antibodies such as hepatitis B immune globulin (HBIG) or botulism immune globulin (BabyBIG®) (63). Hepatitis B immune globulin is commercially available; botulism immune globulin is made

and distributed by the California Department of Public Health and is FDA-approved. A humanized monoclonal antibody recognizing a desired heparin binding motif could be made using technology for the production of palivizumab (Synagis®), a humanized monoclonal antibody that is given monthly to premature newborns to prevent infection with respiratory syncytial virus (64). Because similar heparin binding motifs are also found in *S. epidermidis* and *S. aureus*, using heparin binding motifs as antigens is a first step in developing vaccines against three of the most common causes of central line-associated bloodstream infection.

Antibodies raised against a linear heparin binding motif expressed by a surface protein of *C. albicans* inhibited adhesion of the yeast to heparin in vitro and abolished biofilm formation in vivo. Similar heparin binding motifs occur in surface proteins from *S. epidermidis* and *S. aureus*, two organisms which are also major causes of biofilm-related infections.

In one embodiment, an antibody reactive with a heparin binding motif expressed on the surface of a microorganism is

administered to an individual. The microorganism may be, but is not limited to, a *Candida* species or a *Staphylococcus* species. In one embodiment, the microorganism is *C. albicans*, *S. epidermidis* and/or *S. aureus*. In one embodiment, the heparin binding motif is identified by a computer-based algorithm analysis of protein sequences expressed by a microor-
 5 ganism. In one embodiment, the heparin binding motif to which the antibody was generated is derived from Int1. In one embodiment, the antibody is directed to the peptide HKQEKQKKHQIHKV (SEQ ID NO: 4), or a fragment of
 10 this peptide. In one embodiment, the fragment is at least seven amino acids of the peptide. In one embodiment, the surface expression of the heparin binding motif depends on the life-cycle stage of the microorganism. In one embodiment, the antibody is a polyclonal antibody. In one embodiment, the antibody is a monoclonal antibody.

In one embodiment, a peptide that corresponds to a linear heparin binding motif expressed on the surface of a microor-
 ganism, or a portion of the peptide, is provided. In one
 embodiment, the peptide is used to generate antibodies
 20 directed against the peptide, using methods known in the art. Alternatively, a peptide is provided combined with additional carriers or as a component of a complex, e.g., an adjuvant non-toxic to humans. Such an adjuvant could be aluminum hydroxide used in Engerix-B® (commercially available
 25 hepatitis B vaccine), and in Comvax® (commercially available vaccine for *Haemophilus influenzae* type b and hepatitis B). The peptide corresponds to a linear heparin binding motif, or a portion of the motif, of a microorganism. In one embodi-
 30 ment, the microorganism is a *Candida* species and/or a *Staphylococcus* species. In one embodiment, the microorganism is *C. albicans*, *S. epidermidis* and/or *S. aureus*. In one embodiment, the heparin binding motif is identified by a computer-based algorithm analysis of protein sequences expressed by a microorganism. In one embodiment, the hep-
 35 arin binding motif is derived from Int1. In one embodiment, the peptide is HKQEKQKKHQIHKV (SEQ ID NO: 4), or a fragment of SEQ ID NO: 4. In one embodiment, the peptide is a fragment of the peptide HKQEKQKKHQIHKV (SEQ ID NO: 4), where the fragment contains at least seven amino
 40 acids of the peptide. In one embodiment, the surface expression of the heparin binding motif depends on the life-cycle stage of the microorganism. In one embodiment, the peptide is used to generate an antibody using methods known in the art.

In one embodiment, a method is provided for ameliorating biofilm formation on a surface of an implanted device in a patient. In one embodiment, the method comprises adminis-
 45 tering an antibody to a patient, and ameliorating biofilm formation on a surface of an implanted device in a patient, wherein the antibody is directed to a heparin binding motif expressed on the surface of a microorganism. In one embodi-
 50 ment, the antibody is administered intravenously at a dose not more than 2 grams/kg given as an infusion over 10-24 hours. Since the half-life of antibody preparations is about four weeks, antibody administration would be repeated at monthly
 intervals over the life of the catheter. In one embodiment, the antibody has been modified to increase safety and/or efficacy in a human patient. In one embodiment, the antibody has been humanized using methods known in the art. In one embodi-
 55 ment, the described antibody is provided as a pharmaceutical composition with at least one biocompatible excipient, e.g., buffers, preservatives, tonicity adjusting agents, pH adjusting agents, osmolality adjusting agents, etc., as known to one skilled in the art. Without being held to a single theory, the antibody blocks binding between a heparin binding motif-
 60 containing protein expressed on the surface of a microorgan-

ism and heparin or heparin sulfate, which is expressed by human cells lining the lumen of the implanted device (FIG. 10). Binding of the antibody to the microorganism prevented its attachment to heparin or to heparin sulfate moieties. In the rat model of *Candida* infection of central venous catheters,
 5 the disclosed antibody recognizing a peptide encompassing Motif 1 in *Candida albicans* Int1 (also called anti-KKHQ antibody ("KKHQ" disclosed as SEQ ID NO: 6)) showed excellent in vivo results. The antibody recognizing Motif 1
 10 (anti-KKHQ antibody ("KKHQ" disclosed as SEQ ID NO: 6)) substantially inhibited biofilm production, as shown in FIG. 9. FIGS. 11 and 12A-C demonstrate the anti-KKHQ antibody is effective against various *Candida* species.

The administration of the described antibody is referred to as passive immunization.

In one embodiment, the method comprises administering a peptide to a patient, where the peptide encompasses a heparin binding motif, or a fragment of the motif, that is expressed on
 20 the surface of a microorganism. The peptide is 14 amino acids in length, with amino acids both preceding and following the 8 amino acids of the heparin binding motif. In one embodiment, the peptide is HKQEKQKKHQIHKV (SEQ ID NO: 4), or a fragment thereof. In one embodiment, the described
 25 peptide is provided as a pharmaceutical composition comprising at least one biocompatible excipient including but not limited to buffers, stabilizing agents, solubility enhancing agents, tonicity agents, etc. as known to one skilled in the art.

Without being held to a single theory, the peptide serves as
 30 an antigen to immunize the patient. It provides immunity by promoting endogenous generation of antibodies directed to the peptide, where the endogenously-produced antibodies block binding between heparin, which is administered through the catheter or is bound to a surface of an implanted
 35 medical device, and a heparin binding motif-containing protein expressed on a microorganism surface. Administration of the described peptide is referred to as active immunization. Peptide vaccines are typically administered subcutaneously or intramuscularly. Immunizing dose of the peptide and frequency of immunization are experimentally determined with
 40 humans.

The patient being treated may be naturally immunodeficient by having an underdeveloped immune response, e.g., a
 45 premature newborn. The patient being treated may have a normal immune response, but may be immunocompromised due to particular circumstances, e.g., as a result of treatment such as chemotherapy, or disease, or burns. In one embodi-
 50 ment, the disclosed antibody is administered to an immunodeficient patient. In one embodiment, the disclosed peptide is administered to a non-immunodeficient patient, e.g., an otherwise well adult or child who will receive long term anti-
 biotics through a central line for the treatment of a serious infection, e.g., endocarditis.

In one embodiment, the implanted medical device, e.g., catheter, central line, hemodialysis catheter, peritoneal cath-
 55 eter, PICC line, plastic catheters such as central venous catheters, urinary tract catheters, central nervous system shunt catheters, peritoneal dialysis catheters, dialysis shunts, etc., is implanted in a vein, artery, or a body cavity. In one embodi-
 60 ment, the implanted medical device is a plastic device. In one embodiment, the implanted device is present in the patient's body for a period of time ranging from minutes to six weeks or longer. In one embodiment, the above described methods
 65 may be performed before the device is implanted in the patient. In one embodiment, a first dose of the antibody is administered before the medical device is implanted. In this

embodiment, a peripheral intravenous line could administer this first dose of the antibody if the device to be implanted is a catheter.

In one embodiment, the described methods decrease the virulence of a microorganism, i.e., the method decrease the ability of a microorganism to cause infection in a patient.

All references cited are expressly incorporated by reference herein in their entirety.

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cell line stability, process validation, and scale-up for manufacturing. Biologicals. 1999; 27(3):203-15.

The instant application contains a Sequence Listing which has been submitted in ASCII format via EFS-Web and is hereby incorporated by reference in its entirety. The ASCII copy, created on Mar. 8, 2013, is named 070248_94_SL.txt and is 30,824 bytes in size.

The embodiments described in the specification are only specific embodiments of the inventors who are skilled in the art and are not limiting. Therefore, various changes, modifications, or alterations to those embodiments may be made without departing from the spirit of the invention or the scope of the following claims.

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Asn Thr Asp Leu Phe Lys His Glu Asn Glu Ser Val Asp Leu Leu Leu

-continued

35	40	45
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50	55	60
Ser His Leu Gln Leu Gln	Pro Gln Ser Ser Ser	Ala Ser Ile Phe Asn
65	70	75
Ser Pro Thr Lys	Pro Leu Asn Phe	Pro Arg Thr Asn Ser Lys Pro Ser
	85	90
Leu Asp Pro Asn Ser Ser	Ser Asp Thr Tyr Thr	Ser Glu Gln Asp Gln
	100	105
Glu Lys Gly Lys Glu Glu	Lys Lys Asp Thr Thr	Phe Gln Thr Ser Phe
	115	120
Asp Arg Asn Phe Asp Leu	Asp Asn Ser Ile Asp	Ile Gln Gln Thr Ile
	130	135
Gln His Gln Gln Gln Gln	Pro Gln Gln Gln Gln	Gln Leu Ser Gln Thr
	145	150
Asp Asn Asn Leu Ile Asp	Glu Phe Ser Phe Gln Thr	Pro Met Thr Ser
	165	170
Thr Leu Asp Leu Thr Lys	Gln Asn Pro Thr Val	Asp Lys Val Asn Glu
	180	185
Asn His Ala Pro Thr Tyr	Ile Asn Thr Ser Pro	Asn Lys Ser Ile Met
	195	200
Lys Lys Ala Thr Pro Lys	Val Ser Pro Lys Lys	Val Ala Phe Thr Ala
	210	215
Thr Asn Pro Glu Ile His	His Tyr Pro Asp Asn	Arg Val Glu Glu Glu
	225	230
Asp Gln Ser Gln Gln Lys	Glu Asp Ser Val Glu	Pro Pro Ser Ile Gln
	245	250
His Gln Trp Lys Asp Pro	Ser Gln Phe Asn Tyr	Ser Asp Glu Asp Thr
	260	265
Asn Ala Ser Val Pro Pro	Thr Pro Pro Leu His	Thr Thr Lys Pro Thr
	275	280
Phe Ala Gln Leu Leu Asn	Lys Asn Asn Glu Val	Asn Ser Glu Pro Glu
	290	295
Ala Leu Thr Asp Met Lys	Leu Lys His Glu Asn	Phe Ser Asn Leu Ser
	305	310
Leu Asp Glu Lys Val Asn	Leu Tyr Leu Ser Pro	Thr Asn Asn Asn Asn
	325	330
Ser Lys Asn Val Ser Asp	Met Asp Ser His Leu	Gln Asn Leu Gln Asp
	340	345
Ala Ser Lys Asn Lys Thr	Asn Glu Asn Ile His	Asn Leu Ser Phe Ala
	355	360
Leu Lys Ala Pro Lys Asn	Asp Ile Glu Asn Pro	Leu Asn Ser Leu Thr
	370	375
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Gln Ser Leu Arg Asp Asp	Asn Arg Val Leu Glu	Ser Thr Pro Gly Ser
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Pro Lys Lys Val Asn Pro	Gly Leu Ser Leu Asn	Asp Gly Ile Lys Gly
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Asp Lys Leu Glu Thr Thr	Lys Glu Asn Asp Ala	Pro Glu His Asn Asn
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		460

-continued

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Leu	Val	Ser	Ser	Asp	Asp	His	Leu	Asp	Ser	Phe	Asp	Arg	Ser	Tyr	Asn	485	490	495	
His	Thr	Glu	Gln	Ser	Ile	Leu	Asn	Leu	Leu	Asn	Ser	Ala	Ser	Gln	Ser	500	505	510	
Gln	Ile	Ser	Leu	Asn	Ala	Leu	Glu	Lys	Gln	Lys	Gln	Ile	Gln	Glu	Gln	515	520	525	
Glu	Gln	Thr	Gln	Ala	Ala	Glu	Pro	Glu	Glu	Glu	Thr	Ser	Phe	Ser	Asp	530	535	540	
Asn	Ile	Lys	Val	Lys	Gln	Glu	Pro	Lys	Ser	Asn	Leu	Glu	Phe	Val	Lys	545	550	555	560
Val	Thr	Ile	Lys	Lys	Glu	Pro	Val	Ser	Ala	Thr	Glu	Ile	Lys	Ala	Pro	565	570	575	
Lys	Arg	Glu	Phe	Ser	Ser	Arg	Ile	Leu	Arg	Ile	Lys	Asn	Glu	Asp	Glu	580	585	590	
Ile	Ala	Glu	Pro	Ala	Asp	Ile	His	Pro	Lys	Lys	Glu	Asn	Glu	Ala	Asn	595	600	605	
Ser	His	Val	Glu	Asp	Thr	Asp	Ala	Leu	Leu	Lys	Lys	Ala	Leu	Asn	Asp	610	615	620	
Asp	Glu	Glu	Ser	Asp	Thr	Thr	Gln	Asn	Ser	Thr	Lys	Met	Ser	Ile	Arg	625	630	635	640
Phe	His	Ile	Asp	Ser	Asp	Trp	Lys	Leu	Glu	Asp	Ser	Asn	Asp	Gly	Asp	645	650	655	
Arg	Glu	Asp	Asn	Asp	Asp	Ile	Ser	Arg	Phe	Glu	Lys	Ser	Asp	Ile	Leu	660	665	670	
Asn	Asp	Val	Ser	Gln	Thr	Ser	Asp	Ile	Ile	Gly	Asp	Lys	Tyr	Gly	Asn	675	680	685	
Ser	Ser	Ser	Glu	Ile	Thr	Thr	Lys	Thr	Leu	Ala	Pro	Pro	Arg	Ser	Asp	690	695	700	
Asn	Asn	Asp	Lys	Glu	Asn	Ser	Lys	Ser	Phe	Glu	Asp	Pro	Ala	Asn	Asn	705	710	715	720
Glu	Ser	Ser	Gln	Gln	Gln	Leu	Glu	Val	Pro	His	Thr	Lys	Glu	Asp	Asp	725	730	735	
Ser	Ile	Leu	Ala	Asn	Ser	Ser	Asn	Ile	Ala	Pro	Pro	Glu	Glu	Leu	Thr	740	745	750	
Leu	Pro	Val	Val	Glu	Ala	Asn	Asp	Tyr	Ser	Ser	Phe	Asn	Asp	Val	Thr	755	760	765	
Lys	Thr	Phe	Asp	Ala	Tyr	Ser	Ser	Phe	Glu	Glu	Ser	Leu	Ser	Arg	Glu	770	775	780	
His	Glu	Thr	Asp	Ser	Lys	Pro	Ile	Asn	Phe	Ile	Ser	Ile	Trp	His	Lys	785	790	795	800
Gln	Glu	Lys	Gln	Lys	Lys	His	Gln	Ile	His	Lys	Val	Pro	Thr	Lys	Gln	805	810	815	
Ile	Ile	Ala	Ser	Tyr	Gln	Gln	Tyr	Lys	Asn	Glu	Gln	Glu	Ser	Arg	Val	820	825	830	
Thr	Ser	Asp	Lys	Val	Lys	Ile	Pro	Asn	Ala	Ile	Gln	Ser	Lys	Lys	Phe	835	840	845	
Lys	Glu	Val	Asn	Val	Met	Ser	Arg	Arg	Val	Val	Ser	Pro	Asp	Met	Asp	850	855	860	
Asp	Leu	Asn	Val	Ser	Gln	Phe	Leu	Pro	Glu	Leu	Ser	Glu	Asp	Ser	Gly	865	870	875	880

Phe	Lys	Asp	Leu	Asn	Phe	Ala	Asn	Tyr	Ser	Asn	Asn	Thr	Asn	Arg	Pro	
				885							890			895		
Arg	Ser	Phe	Thr	Pro	Leu	Ser	Thr	Lys	Asn	Val	Leu	Ser	Asn	Ile	Asp	
				900							905			910		
Asn	Asp	Pro	Asn	Val	Val	Glu	Pro	Pro	Glu	Pro	Lys	Ser	Tyr	Ala	Glu	
				915							920			925		
Ile	Arg	Asn	Ala	Arg	Arg	Leu	Ser	Ala	Asn	Lys	Ala	Ala	Pro	Asn	Gln	
				930							935			940		
Ala	Pro	Pro	Leu	Pro	Pro	Gln	Arg	Gln	Pro	Ser	Ser	Thr	Arg	Ser	Asn	
				945							950			955		
Ser	Asn	Lys	Arg	Val	Ser	Arg	Phe	Arg	Val	Pro	Thr	Phe	Glu	Ile	Arg	
				965							970			975		
Arg	Thr	Ser	Ser	Ala	Leu	Ala	Pro	Cys	Asp	Met	Tyr	Asn	Asp	Ile	Phe	
				980							985			990		
Asp	Asp	Phe	Gly	Ala	Gly	Ser	Lys	Pro	Thr	Ile	Lys	Ala	Glu	Gly	Met	
				995							1000			1005		
Lys	Thr	Leu	Pro	Ser	Met	Asp	Lys	Asp	Asp	Val	Lys	Arg	Ile	Leu		
				1010							1015			1020		
Asn	Ala	Lys	Lys	Gly	Val	Thr	Gln	Asp	Glu	Tyr	Ile	Asn	Ala	Lys		
				1025							1030			1035		
Leu	Val	Asp	Gln	Lys	Pro	Lys	Lys	Asn	Ser	Ile	Val	Thr	Asp	Pro		
				1040							1045			1050		
Glu	Asp	Arg	Tyr	Glu	Glu	Leu	Gln	Gln	Thr	Ala	Ser	Ile	His	Asn		
				1055							1060			1065		
Ala	Thr	Ile	Asp	Ser	Ser	Ile	Tyr	Gly	Arg	Pro	Asp	Ser	Ile	Ser		
				1070							1075			1080		
Thr	Asp	Met	Leu	Pro	Tyr	Leu	Ser	Asp	Glu	Leu	Lys	Lys	Pro	Pro		
				1085							1090			1095		
Thr	Ala	Leu	Leu	Ser	Ala	Asp	Arg	Leu	Phe	Met	Glu	Gln	Glu	Val		
				1100							1105			1110		
Arg	His	Pro	Leu	Arg	Ser	Asn	Ser	Val	Leu	Val	His	Pro	Gly	Ala		
				1115							1120			1125		
Gly	Ala	Ala	Thr	Asn	Ser	Ser	Met	Leu	Pro	Glu	Pro	Asp	Phe	Glu		
				1130							1135			1140		
Leu	Ile	Asn	Ser	Pro	Ala	Arg	Asn	Val	Ser	Asn	Asn	Ser	Asp	Asn		
				1145							1150			1155		
Val	Ala	Ile	Ser	Gly	Asn	Ala	Ser	Thr	Ile	Ser	Phe	Asn	Pro	Leu		
				1160							1165			1170		
Asp	Met	Asn	Phe	Asp	Asp	Gln	Ala	Thr	Ile	Gly	Gln	Asn	Ile	Gln		
				1175							1180			1185		
Glu	Gln	Pro	Ala	Ser	Lys	Ser	Ala	Asn	Thr	Val	Arg	Gly	Asp	Asp		
				1190							1195			1200		
Asp	Gly	Leu	Ala	Ser	Ala	Pro	Glu	Thr	Pro	Arg	Thr	Pro	Thr	Lys		
				1205							1210			1215		
Lys	Glu	Ser	Ile	Ser	Ser	Lys	Pro	Ala	Lys	Leu	Ser	Ser	Ala	Ser		
				1220							1225			1230		
Pro	Arg	Lys	Ser	Pro	Ile	Lys	Ile	Gly	Ser	Pro	Val	Arg	Val	Ile		
				1235							1240			1245		
Lys	Lys	Asn	Gly	Ser	Ile	Ala	Gly	Ile	Glu	Pro	Ile	Pro	Lys	Ala		
				1250							1255			1260		
Thr	His	Lys	Pro	Lys	Lys	Ser	Phe	Gln	Gly	Asn	Glu	Ile	Ser	Asn		
				1265							1270			1275		
His	Lys	Val	Arg	Asp												

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1280	1285	1290
His Gln Gln His Asn Pro Ser Met Val Ser Val Pro Ser Gln Tyr		
1295	1300	1305
Thr Asp Ala Thr Ser Thr Asp Pro Asp Glu Asn Lys Asp Val Gln		
1310	1315	1320
His Lys Pro Arg Glu Lys Gln Lys Gln Lys His His His Arg His		
1325	1330	1335
His His His Lys Gln Lys Thr Asp Ile Pro Gly Val Val Asp Asp		
1340	1345	1350
Glu Ile Pro Asp Val Gly Ser Gln Glu Arg Gly Lys Leu Phe Phe		
1355	1360	1365
Arg Val Leu Gly Ile Lys Asn Ile Asn Leu Pro Asp Ile Asn Thr		
1370	1375	1380
His Lys Gly Arg Phe Thr Leu Thr Leu Asp Asn Gly Val His Cys		
1385	1390	1395
Val Thr Thr Pro Glu Tyr Asn Met Asp Asp His Asn Val Ala Ile		
1400	1405	1410
Gly Lys Glu Phe Glu Leu Thr Val Ala Asp Ser Leu Glu Phe Ile		
1415	1420	1425
Leu Thr Leu Lys Ala Ser Tyr Glu Lys Pro Arg Gly Thr Leu Val		
1430	1435	1440
Glu Val Thr Glu Lys Lys Val Val Lys Ser Arg Asn Arg Leu Ser		
1445	1450	1455
Arg Leu Phe Gly Leu Lys Asp Ile Ile Thr Thr Thr Lys Phe Val		
1460	1465	1470
Pro Thr Glu Val Lys Asp Thr Trp Ala Asn Lys Phe Ala Pro Asp		
1475	1480	1485
Gly Ser Phe Ala Arg Cys Tyr Ile Asp Leu Gln Gln Phe Glu Asp		
1490	1495	1500
Gln Ile Thr Gly Lys Ala Ser Gln Phe Asp Leu Asn Cys Phe Asn		
1505	1510	1515
Glu Trp Glu Thr Met Ser Asn Gly Asn Gln Pro Met Lys Arg Gly		
1520	1525	1530
Lys Pro Tyr Lys Ile Ala Gln Leu Glu Val Lys Met Leu Tyr Val		
1535	1540	1545
Pro Arg Ser Asp Pro Arg Glu Ile Leu Pro Thr Ser Ile Arg Ser		
1550	1555	1560
Ala Tyr Glu Ser Ile Asn Glu Leu Asn Asn Glu Gln Asn Asn Tyr		
1565	1570	1575
Phe Glu Gly Tyr Leu His Gln Glu Gly Gly Asp Cys Pro Ile Phe		
1580	1585	1590
Lys Lys Arg Phe Phe Lys Leu Met Gly Thr Ser Leu Leu Ala His		
1595	1600	1605
Ser Glu Ile Ser His Lys Thr Arg Ala Lys Ile Asn Leu Ser Lys		
1610	1615	1620
Val Val Asp Leu Ile Tyr Val Asp Lys Glu Asn Ile Asp Arg Ser		
1625	1630	1635
Asn His Arg Asn Phe Ser Asp Val Leu Leu Leu Asp His Ala Phe		
1640	1645	1650
Lys Ile Lys Phe Ala Asn Gly Glu Leu Ile Asp Phe Cys Ala Pro		
1655	1660	1665
Asn Lys His Glu Met Lys Ile Trp Ile Gln Asn Leu Gln Glu Ile		
1670	1675	1680

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```
Ile Tyr  Arg Asn Arg Phe Arg  Arg Gln Pro Trp Val  Asn Leu Met
    1685                      1690                      1695
```

```
Leu Gln  Gln Gln Gln Gln Gln  Gln Gln Ser Ser Gln  Gln
    1700                      1705                      1710
```

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<210> SEQ ID NO 6
<211> LENGTH: 4
<212> TYPE: PRT
<213> ORGANISM: Candida albicans
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<400> SEQUENCE: 6
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```
Lys Lys His Gln
1
```

```
<210> SEQ ID NO 7
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans
```

```
<400> SEQUENCE: 7
```

```
Ser His Arg Ile Arg Val
1          5
```

```
<210> SEQ ID NO 8
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans
```

```
<400> SEQUENCE: 8
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```
Asn Lys Lys Gly Lys Ser
1          5
```

```
<210> SEQ ID NO 9
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans
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```
<400> SEQUENCE: 9
```

```
Gly Lys Lys Val Lys Ala
1          5
```

```
<210> SEQ ID NO 10
<211> LENGTH: 8
<212> TYPE: PRT
<213> ORGANISM: Candida albicans
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```
<400> SEQUENCE: 10
```

```
Gln Arg Lys Arg Ala Leu Lys Ala
1          5
```

```
<210> SEQ ID NO 11
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans
```

```
<400> SEQUENCE: 11
```

```
Gln Lys Arg Ser Arg Phe
1          5
```

```
<210> SEQ ID NO 12
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans
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<400> SEQUENCE: 12

Asn Lys Lys Phe Lys Asn
1 5

<210> SEQ ID NO 13
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 13

Val Lys Arg Ala Arg Tyr
1 5

<210> SEQ ID NO 14
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 14

Leu Arg Arg Leu Arg Thr
1 5

<210> SEQ ID NO 15
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 15

Ile Lys Lys Asn His Leu
1 5

<210> SEQ ID NO 16
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 16

Gly Lys Lys Val Lys Gln
1 5

<210> SEQ ID NO 17
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 17

Leu Lys Arg Ile His Asn
1 5

<210> SEQ ID NO 18
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 18

Ile Arg Arg Gly Arg Tyr
1 5

<210> SEQ ID NO 19
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 19

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Thr Lys Lys Gly Lys Leu
1 5

<210> SEQ ID NO 20
<211> LENGTH: 8
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 20

Pro His His His Leu Asn Arg Tyr
1 5

<210> SEQ ID NO 21
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 21

Ala Lys Lys Gly Lys Phe
1 5

<210> SEQ ID NO 22
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 22

Thr Lys His Ile His Asn
1 5

<210> SEQ ID NO 23
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 23

Ala Lys Arg Pro Arg Pro
1 5

<210> SEQ ID NO 24
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 24

Leu Arg Lys Tyr Lys Gly
1 5

<210> SEQ ID NO 25
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 25

Gln Arg Lys Leu Lys Thr
1 5

<210> SEQ ID NO 26
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 26

Leu Arg Arg Leu Arg Thr
1 5

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<210> SEQ ID NO 27
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 27

Pro Lys Arg Gln Lys Thr
1 5

<210> SEQ ID NO 28
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 28

Asn Lys His Asn Arg Ile
1 5

<210> SEQ ID NO 29
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 29

Phe His Lys Ala His Gly
1 5

<210> SEQ ID NO 30
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 30

Pro Lys Arg Val Lys Val
1 5

<210> SEQ ID NO 31
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 31

Gly Lys Lys Thr Arg Ile
1 5

<210> SEQ ID NO 32
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 32

Leu Arg Arg Leu Arg Thr
1 5

<210> SEQ ID NO 33
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 33

Pro Lys Lys Leu Lys Phe
1 5

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<210> SEQ ID NO 34
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 34

Gly His Lys Ile Lys Val
1 5

<210> SEQ ID NO 35
<211> LENGTH: 8
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 35

Thr Lys His His Ile Pro Lys Ala
1 5

<210> SEQ ID NO 36
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 36

Ser Lys Arg Asn Lys Asn
1 5

<210> SEQ ID NO 37
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 37

Met Lys Arg Ala Lys Val
1 5

<210> SEQ ID NO 38
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 38

Thr His Arg Phe Lys Phe
1 5

<210> SEQ ID NO 39
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 39

Leu Arg Lys Gly Lys Ser
1 5

<210> SEQ ID NO 40
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 40

Thr His Lys Gly Arg Phe
1 5

<210> SEQ ID NO 41
<211> LENGTH: 6

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<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 41

Ile Lys Lys Phe Lys Tyr
1 5

<210> SEQ ID NO 42
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 42

Cys His Lys Cys His Asn
1 5

<210> SEQ ID NO 43
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 43

Val Lys Lys Phe Arg Gln
1 5

<210> SEQ ID NO 44
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 44

Ala Lys Lys Val Lys Asn
1 5

<210> SEQ ID NO 45
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 45

Val Lys Lys Leu Lys Pro
1 5

<210> SEQ ID NO 46
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 46

Gly Lys Lys Tyr Arg Gly
1 5

<210> SEQ ID NO 47
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 47

Ala Arg Arg Ser Lys Ser
1 5

<210> SEQ ID NO 48
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

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<400> SEQUENCE: 48

Met Lys Arg Gly Lys Pro
1 5<210> SEQ ID NO 49
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 49

Ile Arg His Phe Lys Gly
1 5<210> SEQ ID NO 50
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 50

Phe Lys Arg Phe Lys Phe
1 5<210> SEQ ID NO 51
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 51

Ala His Arg Ala His Ser
1 5<210> SEQ ID NO 52
<211> LENGTH: 8
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 52

Gln Ala Ala His Gln Ile His Lys
1 5<210> SEQ ID NO 53
<211> LENGTH: 8
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 53

Phe Lys Ala Ala Phe Phe Lys Leu
1 5<210> SEQ ID NO 54
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
6xHis tag

<400> SEQUENCE: 54

His His His His His His
1 5<210> SEQ ID NO 55
<211> LENGTH: 35
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 55

gggagctcgt tacttgtcat taattagtta cttec 35

<210> SEQ ID NO 56
<211> LENGTH: 36
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 56

ggacgcggttt ttatcttttt atgtaaatat atacta 36

<210> SEQ ID NO 57
<211> LENGTH: 40
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 57

attgtccaat ttttaaggct gcttttttca aattaatggg 40

<210> SEQ ID NO 58
<211> LENGTH: 40
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 58

ccattaattt gaaaaaagca gccttaaaaa ttggacaatc 40

<210> SEQ ID NO 59
<211> LENGTH: 45
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 59

gcataaacaa gaaaagcagg ccgcccataca aattcataaa gttcc 45

<210> SEQ ID NO 60
<211> LENGTH: 45
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 60

ggaactttat gaatttgatg ggcggcctgc ttttcttggt tatgc 45

<210> SEQ ID NO 61
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence

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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      primer

<400> SEQUENCE: 61

ctcccggcgc ccatggccgc                                20

<210> SEQ ID NO 62
<211> LENGTH: 22
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
      primer

<400> SEQUENCE: 62

gttggtgtgg cccagagact ct                               22

<210> SEQ ID NO 63
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 63

Lys Gln Ala Ala His Gln
1             5

<210> SEQ ID NO 64
<211> LENGTH: 6
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 64

Lys Gln Lys Lys His Gln
1             5

<210> SEQ ID NO 65
<211> LENGTH: 8
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 65

Ile Asp Val Val Asp Gln Ala Lys
1             5

<210> SEQ ID NO 66
<211> LENGTH: 8
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 66

Ser Leu Leu Asp Ala Ala Val Lys
1             5

<210> SEQ ID NO 67
<211> LENGTH: 14
<212> TYPE: PRT
<213> ORGANISM: Candida albicans

<400> SEQUENCE: 67

Val Pro Thr Thr Asp Val Ser Val Val Asp Leu Thr Val Arg
1             5             10

<210> SEQ ID NO 68
<211> LENGTH: 20

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<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 68

tcttgtgaaa ctccgtcgtg 20

<210> SEQ ID NO 69
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 69

agggacgtaa tcaacgcaag 20

<210> SEQ ID NO 70
<211> LENGTH: 19
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 70

tgtgcctggt gcttttacc 19

<210> SEQ ID NO 71
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 71

ttagccaag ctgccattac 20

<210> SEQ ID NO 72
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 72

tcatttgcca ccactaccac 20

<210> SEQ ID NO 73
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 73

tggcatagga ttgtgaccag 20

<210> SEQ ID NO 74
<211> LENGTH: 23
<212> TYPE: DNA

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<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
primer

<400> SEQUENCE: 74

gctggtgggtt attggcaacg tgc 23

<210> SEQ ID NO 75
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
primer

<400> SEQUENCE: 75

tggttaagggtg gtcacggcgg 20

<210> SEQ ID NO 76
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
primer

<400> SEQUENCE: 76

agatcaaggg caaacctcac 20

<210> SEQ ID NO 77
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
primer

<400> SEQUENCE: 77

ataggagcat ttggcacacc 20

<210> SEQ ID NO 78
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
primer

<400> SEQUENCE: 78

taccagggcc aatacaaagg 20

<210> SEQ ID NO 79
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
primer

<400> SEQUENCE: 79

taatgggctt gaccttgag 20

<210> SEQ ID NO 80
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence

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<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 80

ctaattgccgt cgtcagattg 20

<210> SEQ ID NO 81
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 81

aacatctgga acgccatctc 20

<210> SEQ ID NO 82
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 82

ccattgacaa agccggttac 20

<210> SEQ ID NO 83
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 83

ttagatgggt cggattctgg 20

<210> SEQ ID NO 84
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 84

aggtcgcaag caacaacaac 20

<210> SEQ ID NO 85
<211> LENGTH: 21
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 85

aagaaacagc acgagaacca g 21

<210> SEQ ID NO 86
<211> LENGTH: 23
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:

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<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 86

tcctgccact gaaccttccc cag 23

<210> SEQ ID NO 87
 <211> LENGTH: 21
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 87

ccacttgagc cagctggagc g 21

<210> SEQ ID NO 88
 <211> LENGTH: 20
 <212> TYPE: DNA
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 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 88

ccacccaaac caagtgcctac 20

<210> SEQ ID NO 89
 <211> LENGTH: 20
 <212> TYPE: DNA
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 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 89

aactccagat gatcccgaag 20

<210> SEQ ID NO 90
 <211> LENGTH: 20
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 90

tgtgcccact gaagtcaaag 20

<210> SEQ ID NO 91
 <211> LENGTH: 20
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic primer

<400> SEQUENCE: 91

gctttaccgg tgatttggtc 20

<210> SEQ ID NO 92
 <211> LENGTH: 20
 <212> TYPE: DNA
 <213> ORGANISM: Artificial Sequence
 <220> FEATURE:
 <223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic

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    primer

<400> SEQUENCE: 92

cacctcatgc tccaacaatg                                20

<210> SEQ ID NO 93
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
    primer

<400> SEQUENCE: 93

gatgattctg gggctgattc                                20

<210> SEQ ID NO 94
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
    primer

<400> SEQUENCE: 94

tgctgaaagt tctgcaccag                                20

<210> SEQ ID NO 95
<211> LENGTH: 20
<212> TYPE: DNA
<213> ORGANISM: Artificial Sequence
<220> FEATURE:
<223> OTHER INFORMATION: Description of Artificial Sequence: Synthetic
    primer

<400> SEQUENCE: 95

gcttcaacgg aaacagaagc                                20

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What is claimed is:

1. A composition comprising at least one biocompatible excipient and an antibody capable of binding to peptide HKQEKQKKHQIHKV (SEQ ID NO: 4) wherein the excipient comprises a Tris buffer and wherein the antibody is a non-human humanized antibody.

2. The composition of claim 1, where the peptide HKQEKQKKHQIHKV (SEQ ID NO: 4) is a portion of a protein.

3. The composition of claim 2 where the microorganism is a *Candida* species or a *Staphylococcus* species.

4. The composition of claim 1 where the peptide is derived from Int1 protein (SEQ ID NO: 5).

5. The composition of claim 1 where the antibody is a polyclonal antibody.

6. The composition of claim 1 where the antibody is a monoclonal antibody.

7. A kit comprising instructions for administering a composition to a patient candidate for medical device installation or implantation, and

a composition comprising at least one biocompatible excipient and an antibody capable of binding to peptide HKQEKQKKHQIHKV (SEQ ID NO: 4) where the excipient comprises a Tris buffer and wherein the antibody is a non-human humanized antibody.

8. An antibody capable of binding to peptide HKQEKQKKHQIHKV (SEQ ID NO: 4) where the antibody is a nonhuman humanized antibody.

* * * * *